

# METHOD FOR DETERMINING AND CORRECTING VISION

## Cross Reference To Related Applications

This application is a continuation of application Serial No. 09/566,668 filed May 8, 2000 for "Apparatus and Method for Objective Measurement and Correction of Optical Systems," which itself is a continuation-in-part of application Serial Number 09/324,179 filed May 20, 1998 for "Objective Measurement and Correction of Optical Systems Using Wavefront Analysis," which itself is a continuation of application Serial Number 08/756,272 filed November 25, 1996 for "Objective Measurement and Correction of Optical Systems Using Wavefront Analysis," now abandoned, all of which are commonly owned and have the disclosures incorporated by reference.

## Field of the Invention

The invention relates generally to optical aberration measurement and correction, and more particularly to an objective measurement and correction of optical systems, such as systems of a human eye.

## Background of the Invention

Optical systems having a real image focus can receive collimated light and focus it at a point. Such optical systems can be found in nature, e.g., human and animal eyes, or can be man-made, e.g., laboratory systems, guidance systems, and the like. In either case, aberrations in the optical system can affect the system's performance. By way of example, the human eye will be used to explain this problem.

A perfect or ideal eye diffusely reflects an impinging light beam from its retina through optics of the eye which includes a lens and a cornea. For such an ideal eye in a relaxed state, i.e., not accommodating to provide near-field focus, reflected light exits the eye as a sequence of plane waves. However, an eye typically has aberrations that cause deformation or distortion of reflected light waves exiting the eye. An aberrated eye diffusely reflects an impinging light beam from its retina through its lens and cornea as a sequence of distorted wavefronts.

There are a number of technologies that attempt to provide the patient with improved visual acuity. Examples of such technologies include remodeling of the cornea using refractive laser surgery or intra-corneal implants, adding synthetic lenses to the optical system using intra-ocular lens implants, and precision-ground spectacles.

5 In each case, the amount of corrective treatment is typically determined by placing spherical and/or cylindrical lenses of known refractive power at the spectacle plane (approximately 1.0-1.5 centimeters anterior to cornea) and literally asking the patient which lens or lens combination provides the clearest vision. This is an imprecise measurement of true distortions in the reflected wavefront because 1) a single sphero-

10 cylindrical compensation is applied across the entire wavefront, 2) vision is tested at discrete intervals (i.e., diopter units) of refractive correction, and 3) subjective determination by the patient is desired in order to determine the optical correction. Thus, conventional methodology for determining refractive errors in the eye is substantially less accurate than the techniques now available for correcting the ocular aberrations.

15

One method of measuring ocular refractive errors is disclosed in U.S. Patent No. 5,258,791 to Penney et al. for "Spatially Resolved Objective Autorefractometer," which teaches the use of an autorefractometer to measure the refraction of the eye at numerous discrete locations across the corneal surface. The autorefractometer is

20 designed to deliver a narrow beam of optical radiation to the surface of the eye, and to determine where that beam strikes the retina using a retinal imaging system. Both the angle of the beam's propagation direction with respect to the optical axis of the system and the approximate location at which the beam strikes the corneal surface of the eye are independently adjustable. However, a small uncertainty or error in the location of the beam's point of incidence on the cornea exists due to the curved corneal surface.

25 For each point of incidence across the corneal surface, the refraction of the eye corresponding to that surface point can be determined by adjusting the angle at which the beam strikes the cornea until the beam refracted on to the iris strikes the fovea centralis. Adjustment of the beam angle of propagation can be accomplished either

manually by the patient or automatically by the autorefractometer, if a feedback loop involving a retinal imaging component is incorporated.

Penney '791 further teaches the use of the autorefractometer measurements in determining the appropriate corneal surface reshaping to provide emmetropia, a condition of a normal eye when parallel beams or rays of light are focused exactly on the retina and vision is perfect. This is accomplished by first obtaining an accurate measurement of corneal surface topography using a separate commercially available device. A mathematical analysis is then performed using an initial corneal topography at each surface reference point, the measured refraction at each surface point, and Snell's law of refraction, to determine a desired change in surface contour at each reference point. The contour changes at the various reference points are then combined to arrive at a single reshaping profile to be applied across the full corneal surface.

A major limitation to the approach described by Penney '791 is that a separate measurement of corneal topography is desired to perform the Snell's Law analysis of needed refraction change. This adds significantly to the time and cost of a complete and desirable diagnostic evaluation. Further, the accuracy of the refraction change analysis will be dependent upon the accuracy of the topographic measurement and the accuracy of the autorefractometer measurement. In addition, any error in the spatial orientation of a topography map with respect to a refraction map will degrade the accuracy of the needed correction profile. Yet another limitation to known approaches such as described in Penney '791, by way of example, is that test points on the corneal surface are examined sequentially. Eye motion during the examination, either voluntary or involuntary, could introduce substantial errors in the refraction measurement. Penney '791 teaches detection of such eye movement by deliberately including measurement points outside the pupil, i.e., in the corneal region overlying the iris, where the return from the retina will obviously be zero at specific intervals in the examination sequence. However, this approach may still allow substantial undetected eye movement error between such iris reference points.

By way of example, one method and system known in the art, are disclosed by Junzhong Liang et al. in "Objective Measurement Of Wave Aberrations Of The Human Eye With The Use Of A Hartmann-Shack Wave-Front Sensor," published in the Journal of the Optical Society of America, Volume 11, No. 7, July 1994, pages 1949-1957. Liang et al. teach the use of a Hartmann-Shack wavefront sensor to measure ocular aberrations by measuring the wavefront emerging from the eye by the retinal reflection of a focused laser light spot on the retina's fovea. The actual wavefront is reconstructed using wavefront-estimation with Zernike polynomials.

The imprecise measurement technique of placing lenses of known refractive power anterior to the cornea and asking a patient which lens or lens combination provides the clearest vision has been improved with the use of autorefractometers, as described in Penny '79, or with the use of wavefront sensors as described by Liang et al. Spatially resolved refraction data, in combination with measured existing surface contour of the anterior surface of the eye, enable a calculation of a detailed spatially resolved new contour which provides corrected vision. However, it would be an improvement in this art if such vision correction could be made without the need for this contour data, and further without the need for feedback from the patient regarding an appropriate lens. Liang et al. discloses the use of a Hartmann-Shack wavefront sensor to measure ocular aberrations by measuring the wavefront emerging from the eye by retinal reflection of a focused laser light spot on the retina's fovea. A parallel beam of laser light passes through beam splitters and a lens pair which brings the beam to a focus point on the retina by the optics of the eye. Possible myopia or hyperopia of the tested eye is corrected by movement of a lens within the lens pair. The focused light on the fovea is then assumed to be diffusely reflected and acts as a point source located on the retina. The reflected light passes through the eye and forms a distorted wavefront in front of the eye that results from the ocular aberrations. The aberrated wavefront is then directed to the wavefront sensor.

A point source of radiation on the retina would be ideal for such measurements. However, when the perfect eye receives a collimated beam of light, the best possible image on the retina is a diffraction limited spot. As illustrated by way of example, with

Penny et al. and Liang et al., discussed above, and typical for those of skill in the art, parallel or collimated beams are used with the optics of the eye being measured to achieve this diffraction limited spot for such objective measurements. To do so, a setup for each patient includes a corrective lens or lens combination and adjustments thereto for accommodating that patient's specific visual acuity. Providing a corrective or lens combination, as well as setting up for their use becomes cumbersome, time consuming, and at an additional expense. Eliminating the need for such corrective optics is desirable and eliminates a variable within optical measurement systems that typically include many variables. Further, there is a need for providing optical characteristics of an eye without requiring feedback from the patient. By way of example, the patient may be a wild or domestic animal, living or dead.

The Hartmann-Shack wavefront sensor disclosed by Liang et al. includes two identical layers of cylindrical lenses with the layers arranged so that lenses in each layer are perpendicular to one another, as further disclosed in U.S. Patent No. 5,062,702 to Bille. In this way, the two layers operate as a two-dimensional array of spherical lenslets that divide the incoming light wave into sub-apertures. The light through each sub-aperture is brought to focus in the focal plane of the lens array where a charge coupled device (CCD) image module resides.

The system of Liang et al. is calibrated by impinging an ideal plane wave of light on the lenslet array so that a reference or calibrating pattern of focus spots is imaged on the CCD. Since the ideal wavefront is planar, each spot related to the ideal wavefront is located on the optical axis of the corresponding lenslet. When a distorted wavefront passes through the lenslet array, the image spots on the CCD are shifted with respect to a reference pattern generated by the ideal wavefront. Each shift is proportional a local slope, i.e., partial derivatives of the distorted wavefront, which partial derivatives are used to reconstruct the distorted wavefront, by means of modal wavefront estimation using Zernike polynomials.

However, the system disclosed by Liang et al. is effective only for eyes having fairly good vision. Eyes that exhibit considerable myopia (near-sightedness) would cause the focus spots to overlap on the CCD, thereby making local slope determination

practically impossible for eyes having this condition. Similarly, eyes that exhibit considerable hyperopia (farsightedness) deflect the focus spots such that they do not impinge on the CCD thereby again making local slope determination practically impossible for eyes having this condition.

5

### **Summary of the Invention**

In general, an embodiment of the present invention provides a method and system for objectively measuring aberrations of optical systems by wavefront analysis and use such measurement to generate an optical correction. Another embodiment further provides for the objective measurement of ocular aberrations having a dynamic range that can cope with large amounts of such aberrations so as to be useful in practical applications. Still another embodiment of the present invention provides a method and system for objectively measuring ocular aberrations using a wavefront analyzer of simple and inexpensive design.

10

15

20

25

One embodiment of the present invention provides an apparatus and method for making objective and detailed measurements of aberrations present in human eyes. Aberrations measured by the apparatus include "higher order" phenomena, such as spherical aberration and coma, in addition to the traditional myopia/hyperopia and astigmatism. Once the apparatus obtains data representing aberration information, this data is transferred to a treatment system which may employ a small diameter treatment laser beam, may employ a computer controlled laser pulse placement, and may employ an active eye-tracking module. These treatment system features permit corrective laser surgery to address, and ideally to eliminate, the aberrations measured by the apparatus. Another means of correction may be employed, such as an embodiment of the present of the present invention which improves visual performance of treated eyes beyond the level obtained by current refractive procedures.

30

In accordance with an embodiment of the present invention, an energy source generates a beam of radiation. Optics, disposed in the path of the beam, direct the beam through a focusing optical system that has a rear portion which provides a diffuse reflector. The beam is diffusely reflected back from the rear portion as a wavefront of

radiation that passes through the focusing optical system to impinge on the optics. The optics project the wavefront to a wavefront analyzer in direct correspondence with the wavefront as it emerges from the focusing optical system. A wavefront analyzer is disposed in the path of the wavefront projected from the optics and calculates distortions of the wavefront as an estimate of ocular aberrations of the focusing optical system. The wavefront analyzer includes a wavefront sensor coupled to a processor that analyzes the sensor data to reconstruct the wavefront to include the distortions thereof.

One embodiment of the present invention, herein described by way of example, utilizes wavefront sensing to measure the aberrations of the eye. When one considers the perfect or ideal eye as earlier described, a perfectly collimated light beam (i.e., a bundle of parallel light rays) incident on the perfect, ideal emmetropic eye, focuses to a diffraction-limited small spot on the retina. This perfect focusing is true for all light rays passing through the entrance pupil, regardless of position. From the wavefront perspective, the collimated light represents a series of perfect plane waves striking the eye. Due to the reversible nature of light ray propagation, the light emanates from an illuminated spot created on the retina as wavefronts exiting the ideal eye as a series of perfect plane waves. The apparatus of the present invention achieves this ray reversal effect using a probe beam optical path for projecting a small diameter, eye-safe laser beam into the eye and onto the fovea. The light scattered from the irradiated retina serves as a secondary source for a re-emitted wavefront. The probe laser beam strikes the retina at an appropriate foveal location to illuminate a sufficiently small spot. A fixation optical path is provided which includes a reference target aligned to an optical axis. This allows a patient to fixate on a target. A video path provides a video image of the eye plane, centered on the optical axis. A video image of the eye allows a clinical operator to assist in orienting the eye for the wavefront measurement.

Embodiments of the present invention, herein described, provide a refraction measurement system that easily accommodates the measurement of vision characteristics of the eye, even in the presence of finite refractive errors. The time for a patient to be in a fixed position during examination is reduced, while at the same time

providing a useful source of light on the retina of the eye to be measured regardless of the characteristics of the eye of that patient or other patients to be examined.

Desirably, measurements are made without requiring patient or operator feedback.

One method aspect of the invention for measuring optical characteristics of an optical system, such as the eye, includes focusing an optical beam onto an anterior surface of the eye for providing a finite source of secondary radiation on the retina of the eye, which secondary radiation is emitted from the retina as a reflected wavefront of radiation that passes through the eye. The reflected wavefront is directed onto a wavefront analyzer for measuring distortions associated with the reflected wavefront.

One method aspect of the present invention includes a method for enhancing vision in an eye, which method comprises determining an optical path difference between a plane wave and a wavefront emanating from a region of the retina of the eye, and optically correcting for visual defects of the eye based on the optical path difference and refractive indices of media through which the wave front passes, to thereby cause the wavefront to approximate the shape of the plane wave. One embodiment herein described includes an apparatus having an optical correction system comprising a wavefront analyzer disposed in the path of a wavefront emanating from the eye for determining an optical path difference between a plane wave and the wavefront, and a converter for providing an optical correction based on the path difference and refractive indices of media through which the wavefront passes. Such an embodiment of the present invention enables treatment of the eye to permit each treated eye to function just as an ideal emmetropic eye. With a difference between a complex reflected wavefront and an ideal plane wave, an optical path difference (OPD) exists at each transverse location of the wavefronts. Consider a light ray propagating through the eye and intersecting the cornea at some location  $(x, y)$ . Laser ablation to a depth  $d$  at that site reduces the optical path difference by the amount  $(n-n_0)d$ , where  $n$  is the index of refraction of corneal tissue, and  $n_0$  is equal to 1, the index of refraction of air. The entire aberrated wavefront is corrected by measuring the OPD at each  $(x, y)$  location and ablating the cornea to a depth profile  $d(x, y)$  so that:  $d(x,y) = \text{OPD}(x,y) \div (n-1)$ . Thus, the optimal ablation profile for correction of the measured aberrations is



essentially the OPD profile scaled by the refractive index difference. An embodiment of the invention measures the shape of the re-emitted wavefront, so that an appropriate amount of treatment laser exposure for each corneal location can then be calculated from the optimal ablation profile, along with factors such as the spatial effectiveness of the laser ablation as a function of corneal position.

In one embodiment, the radiation is optical radiation and the wavefront sensor is implemented using a plate and a planar array of light-sensitive cells. The plate is generally opaque but that has an array of light transmissive apertures that selectively let impinging light therethrough. The plate is disposed in the path of the wavefront so that portions of the wavefront pass through the light transmissive apertures. The planar array of cells is arranged parallel to and spaced apart from the plate by a selected distance. Each portion of the wavefront passing through one of the light transmissive apertures illuminates a geometric shape covering a unique plurality of cells.

As herein described, by way of example, the wavefront optical path of the present invention relays the re-emitted wavefront from the corneal plane to an entrance face of a Hartman-Shack wavefront sensor. The wavefront incident on the sensor is received by a sensitive charged-coupled device (CCD) camera and an optical plate containing an array of lenslets. The lenslet array is parallel to the CCD detector face with a distance therebetween approximately equal to the focal length of each lens in the lenslet array. The lenslet array divides the incoming wavefront into a matching array of "wavelets," each of which focuses to a small spot on the CCD detector plane. The constellation of wavelet spots in the CCD is used to reconstruct the shape of the incident wavefront. Collimated light striking the lenslet at normal (perpendicular) incidence would focus to the spot on the CCD face where this optical axis intersects. The optics of the apparatus provides such collimated light to the wavefront sensor using a calibration optical path. Collimated light CCD images are routinely obtained as part of a daily calibration process and used for reference in analyzing experimental data.

However, in the case of a reflected aberrated wave front, light focuses to a spot displaced from the collimated reference point by a distance  $D_x$ . The distance from the lenslet face to the CCD surface,  $D_z$ , is precisely known. Therefore, dividing the

measured displacement,  $Dx$ , by the known propagation distance,  $Dz$ , the slope of the wavefront at the location of this lens element is determined. The same calculation is applied in the  $y$  direction within the plane, and the entire process applied to every lenslet element irradiated by the wavefront. A mathematical algorithm is then applied to reconstruct the wavefront shape consistent with the calculated  $Dx/Dz$  and  $Dy/Dz$  slope data. Regardless of which wavefront sensor is used, the distance between the planar array of cells and the opaque plate, or the array of lenslets, can be varied to adjust the slope measurement gain of the wavefront sensor and thereby improve the dynamic range of the system.

Another measure of dynamic range enhancement is provided by the focusing optics. The focusing optics includes first and second lenses maintained in fixed positions in the path of the beam and wavefront. An arrangement of optical elements is disposed between the lenses in the path of the beam and the wavefront. The optical elements are adjustable to change the optical path length between the lenses. If an optical correction is desired, the distortions are converted to an optical correction which, if placed in the path of the wavefront, causes the wavefront to appear approximately as a plane wave. The optical correction can be in the form of a lens or an amount of corneal material ablated from the eye.

An embodiment of the present invention provides a method for enhancing vision in an eye, the method comprising determining an optical path difference between a plane wave and a wavefront emanating from an eye, producing a plurality of laser beam shots, applying said plurality of laser beam shots to the eye in a manner that is based in part on the optical path difference between the plane wave and the wavefront emanating from the eye, and removing tissue from the cornea of the eye in a manner that reduces the optical path difference between the plane wave and the wavefront emanating from the eye whereby visual defects of the eye are reduced. Further embodiments of this embodiment provide that the size of a laser beam shot is less than about 1 mm, is less than about 0.5 mm, or that the size of the laser beam shot varies.

An embodiment of the present invention provides a method for enhancing vision in an eye requiring a myopic correction of greater than  $-3$  diopters to an eye having

perfect vision, a myopic correction of greater than  $-3$  diopters to an eye having about 20/20 vision, a myopic correction of greater than  $-3$  diopters to an eye having better than 20/20 vision, a myopic correction of greater than  $-3$  diopters to an eye having at least 20/10 vision, a myopic correction of greater than  $-6$  diopters to an eye having perfect vision, a myopic correction of greater than  $-6$  diopters to an eye having about 20/20 vision, a myopic correction of greater than  $-6$  diopters to an eye having better than 20/20 vision, a myopic correction of greater than  $-6$  diopters to an eye having at least 20/10 vision, a myopic correction of greater than  $-8$  diopters to an eye having perfect vision, a myopic correction of greater than  $-8$  diopters to an eye having about 20/40 vision, a myopic correction of greater than  $-8$  diopters to an eye having better than 20/40 vision, a myopic correction of greater than  $-8$  diopters to an eye having at least 20/20 vision, a hyperopic correction of greater than  $+3$  diopters to an eye having perfect vision, a hyperopic correction of greater than  $+3$  diopters to an eye having about 20/20 vision, a hyperopic correction of greater than  $+3$  diopters to an eye having better than 20/20 vision, a hyperopic correction of greater than  $+3$  diopters to an eye having at least 20/10 vision, a hyperopic correction of greater than  $+6$  diopters to an eye having perfect vision, a hyperopic correction of greater than  $+6$  diopters to an eye having about 20/20 vision, a hyperopic correction of greater than  $+6$  diopters to an eye having better than 20/20 vision, a hyperopic correction of greater than  $+6$  diopters to an eye having at least 20/10 vision, a hyperopic correction of greater than  $+8$  diopters to an eye having perfect vision, a hyperopic correction of greater than  $+8$  diopters to an eye having about 20/40 vision, a hyperopic correction of greater than  $+8$  diopters to an eye having better than 20/40 vision, or a hyperopic correction of greater than  $+8$  diopters to an eye having at least 20/20 vision. The method comprises determining an optical path difference between a plane wave and a wavefront emanating from an eye, producing a plurality of laser beam shots, applying said plurality of laser beam shots to the eye in a manner that is based in part on the optical path difference between the plane wave and the wavefront emanating from the eye, and removing tissue from the cornea of the eye in a manner that reduces the optical path difference between the plane wave and the wavefront emanating from the eye whereby visual defects of the

eye are reduced. Further embodiments of this embodiment provide that the size of a laser beam shot is less than about 1 mm, is less than about 0.5 mm, or that the size of the laser beam shot varies.

5 An embodiment of the present invention provides a method for enhancing vision in an eye, the method comprising determining an optical path difference between a plane wave and a wavefront emanating from an eye, producing a plurality of laser beam shots; mechanically removing the epithelium of the eye to expose Bowmans membrane; applying said plurality of laser beam shots to the Bowmans membrane in a manner that is based in part on the optical path difference between the plane wave and the  
10 wavefront emanating from the eye, and said plurality of laser beam shots removing tissue from the eye in a manner that reduces the optical path difference between the plane wave and the wavefront emanating from the eye, whereby the vision of the eye is improved.

15 An embodiment of the present invention provides a method for enhancing vision in an eye, the method comprising, determining an optical path difference between a plane wave and a wavefront emanating from an eye, producing a plurality of laser beam shots, displacing a portion of the eye to expose the stroma of the eye, such as by way of example using a lasik procedure or cutting and removing a lenticle from the anterior surface of the cornea, applying said plurality of laser beam shots to the exposed stroma  
20 in a manner that is based in part on the optical path difference between the plane wave and the wavefront emanating from the eye, said plurality of laser beam shots removing tissue from the eye in a manner that reduces the optical path difference between the plane wave and the wavefront emanating from the eye, and replacing the displaced portion of the eye; whereby the vision of the eye is improved.

25 A further embodiment of the present invention provides a method for enhancing vision in an eye, the method comprising, determining an optical path difference between a plane wave and a wavefront emanating from an eye, producing a plurality of laser beam shots, applying said plurality of laser beam shots to the eye in a manner to create two different focus zones and that is based in part on the optical path difference  
30 between the plane wave and the wavefront emanating from the eye, and said plurality

of laser beam shots removing tissue from the eye in a manner that reduces the optical path difference between the plane wave and the wavefront emanating from the eye; whereby the vision of the eye is improved.

A method aspect of the present invention, as herein described, determines  
5 aberrations of an eye requiring greater than a + or - 3 diopter correction, and includes directing an optical beam onto a retina of an eye, reflecting the optical beam from the retina of the eye, determining characteristics of a wavefront in a reflected optical beam, and generating data based on the characteristics of the wavefront, which data  
10 quantifies the aberrations of the eye. The data may further be generated based on refractive indices of media through which the optical beam passes. Yet further, data based on the characteristics of the wavefront, which data quantifies the aberrations of the eye for a discrete section of the eye may also be generated.

One method for determining aberrations of an eye, herein described by way of  
15 example, includes directing a probe beam along a probe beam path toward an eye, directing a fixation image along a fixation image path toward the eye, directing a light source along a video image path toward the eye, generating a video image of the eye, directing a wavefront originating from the eye along a wavefront path, wherein the probe beam path, the fixation image path, the video image path, and the wavefront path  
20 are coincident at least along a portion of their respective paths, the probe beam path terminating at the retina of the eye and the probe beam reflecting from the retina of the eye as a wavefront, aligning the eye with the probe beam path based at least in part on the video image of the eye generated by the light source directed along the video image path, measuring the wavefront, and generating data representative of the aberrations of the eye based on the measurement of the wavefront. Further, the aligning of the eye  
25 with the probe beam path based at least in part on the video image of the eye generated by the light source directed along the video image path, may have the wavefront pass through a single microlens array.

One apparatus for determining the aberrations of an eye comprises a patient  
30 head rest comprising vertical adjustment, the patient head rest associated with an optical table having a base. The base carries a probe beam generating apparatus,

probe beam directing optics, the probe beam directing optics comprising a beam splitter; a mirror; and a lens, the probe beam directing optics being capable of directing a probe beam toward an eye of a patient positioned on the patient head rest, video image components, the video image components comprising a light source, a mirror, and a video camera, the video image components being capable of generating an image of an eye of a patient positioned on the patient head rest, eye fixation components, the eye fixation component comprising a fixation target; a light source; a lens; and a mirror, the fixation components being capable of generating a target that the eye of a patient positioned on the patient head rest can view, and wavefront directing and analyzing components, the wavefront directing and analyzing components comprising a lens, a mirror, a microlens array, a camera, and a data processor. The wavefront directing and analyzing components are capable of measuring the wavefront emanating from the eye of a patient positioned on the patient head rest and determining aberrations of said eye that range from at least about + or - 1 diopters to at least about + or - 6 diopters.

### **Brief Description of the Drawings**

Embodiments of the invention are described by way of example with reference to the accompanying drawings in which:

FIG. 1A is a schematic view of the ideal eye reflecting light from its retina as a planar wavefront;

FIG. 1B is a schematic view of an aberrated eye reflecting light from its retina as a deformed wavefront;

FIG. 1C is a schematic view of the distorted wavefront relative to a reference plane to show the wavefront error or optical path difference as a function of transverse distance in the propagation direction;

FIG. 1D is a schematic view illustrating use of a reference plane;

FIG. 2 is a simplified schematic of the system for determining ocular aberrations in accordance with the essential features of the present invention;

FIG. 3 is a schematic of one embodiment of a Hartmann-Shack wavefront analyzer used in the present invention;

FIG. 4 is a perspective view of a portion of the pinhole imaging plate and planar array of light-sensitive cells comprising the wavefront sensor from the embodiment of FIG. 3 where the deflection of a wavefront piece associated with an aberrated eye is shown in comparison with a wavefront piece associated with a calibration or planar wavefront;

FIG. 5 is a plan view of a designated area on the planar array of light-sensitive cells associated with a corresponding hole;

FIG. 6 is a schematic of another embodiment of a wavefront analyzer used in the present invention;

FIG. 7 is a schematic view of an embodiment of the present invention suitable for ophthalmic use;

FIG. 8 is a side view of a cornea showing a thickness of corneal material to be ablated as an optical correction generated by the present invention;

FIG. 9 is a side elevation view of one embodiment of the present invention illustrating a patient positioning for measurement;

FIG. 10 is an end elevation view of the embodiment of FIG. 9;

FIG. 11 is an enlarged perspective view of an patient positioning portion of the embodiment of FIG. 9;

FIG. 12 is a top plan view of optical elements of the embodiment of FIG. 9;

FIG. 12A illustrates a fixation target optical path of FIG. 12;

FIG. 12B illustrates a video image optical path of FIG. 12;

FIG. 12C illustrates a probe laser optical path of FIG. 12;

FIG. 12D illustrates a re-emitted wavefront optical path of FIG. 12;

FIG. 12E illustrates a calibration wavefront optical path of FIG. 12;

FIGS. 12F and 12G are front elevation and top plan views of a trial lens holder useful with embodiments of the present invention herein described;

FIG. 13 is a block diagram illustrating electrical components of the embodiment of FIG. 9;

FIG. 14 is an enlarged image of an eye illustrating a centration image;

FIG. 15 is a block diagram illustrating an operable flow of steps used in one embodiment of the present invention;

FIG. 16 is an enlarged image of an eye illustrating a pre-measurement eye alignment;

FIG. 17 is an enlarged image of an eye illustrating a pre-measurement eye alignment checking thereof;

FIG. 18 is a line diagram illustrating an eye registration pattern;

FIG. 19 illustrates a rejected CCD image;

FIG. 20 illustrates a CCD image including centroids;

FIG. 21 is an enlarged image of a centroid;

FIG. 22 illustrates an image available to an operator of a measured and reference centroid;

FIG. 23A illustrates a spacial filter operable in one embodiment of the present invention;

FIG. 23B illustrates a noisy CCD image before filtering to provide an image as illustrated with reference to FIG. 20;



FIG. 24A is a three dimensional plot of a wavefront reconstruction in accordance with the present invention;

FIG. 24B illustrates a higher order aberration for the wavefront of FIG. 23;

5      FIG. 25 illustrates a geometric effect of a curved corneal surface on a wavefront measurement;

FIGS. 26A and 26B illustrate ablation depth profiles for surgery on a myopic eye and a hyperopic eye, respectively;

FIG. 26C illustrates an ablation efficiency function for one embodiment of the present invention;

10      FIG. 27A is a pictorial line drawing illustrating magnification modification to the embodiment of FIG. 12; and

FIG. 27B is a pictorial line drawing illustrating optical elements of the present invention.

### **Detailed Description of the Invention**

The present invention will now be described more fully hereinafter with reference to the accompanying drawings, in which embodiments of the present invention are shown by way of illustration and example. This invention may, however, be embodied in many forms and should not be construed as limited to the embodiments set forth herein. Rather, these embodiments are provided so that this disclosure will be thorough and complete, and will fully convey the scope of the invention to those skilled in the art. Like numbers refer to like elements throughout.

By way of illustrative example, the present invention will be described with respect to diagnosing and correcting a human eye. However, it is to be understood that the teachings of the present invention are applicable to any optical system having a real image focus that can be, or can be adapted to diffusely reflect a focused spot of radiation from a rear portion of the optical system back through the optical system as a wavefront of radiation. Thus, the present invention can be used with human or animal eyes of patients that may be alive or dead, or any man-made optical system.

Correction of the human eye that may be used in conjunction with or based upon the diagnostic information provided by embodiments of the present invention include, by way of example, the grinding or preparation of eye glasses and lenses, which teachings are well known in the art, such as described in "Geometric, Physical, and Visual Optics" by Michael P. Keating, Ph.D. published by Butterworth Publishers, 80 Montvale Avenue, Stone, MA 02180, Copyright 1988, herein incorporated by reference. Laser surgery using lasers that photo ablate corneal tissue through the use of broad beam excimer lasers which are well known in the art, such as those disclosed in U.S. Patent No. 5,163,934 to Trokel, correction of presbyopia by photorefractive keratectomy disclosed in U.S. Patent No. 5,395,356 to King et al., and narrow beam systems as described in U.S. Patent No. 5,849,006 to Frey et al. in conjunction with a Lasik procedure which are well known in the art, the disclosures of which are herein incorporated by reference.

The method of using wavefront analysis to determine an appropriate optical correction will be introduced with reference to the eye example and the aid of the

schematic drawings of FIGS. 1A, 1B, and 1C. As earlier described with reference to an ideal eye, and with reference now to FIG. 1A, the ideal emmetropic or perfect eye **100** diffusely reflects an impinging light beam (not shown for sake of clarity) from the back of its retina **102** (i.e., the fovea centralis **103**) through the eye's optics which includes lens **104** and cornea **106**. For such an ideal eye **100** in a relaxed state, i.e., not accommodating to provide near-field focus, the reflected light (represented by arrows **108**) exits the eye **100** as a sequence of plane waves, one of which is represented by straight line **110**. However, as illustrated with reference to FIG. 1B, a typical eye **120** normally has aberrations that cause deformation or distortion of a reflected wave exiting the eye, where the aberrated eye **120** diffusely reflects an impinging light beam (again not shown for sake of clarity) from the back of its retina **122** of the fovea centralis **123** through lens **124** and cornea **126**. For the aberrated eye **120**, the reflected light **128** exits the eye **120** as a sequence of distorted wavefronts, one of which is represented by wavy line **130**.

With reference now to FIG. 1C, a coordinate system is defined for convenience, where positive x is upward in the plane of the figure, positive y is outward from the plane of the figure, and positive z is to the right along a propagation direction. The distorted wavefront **130** is herein described mathematically as  $W(x,y)$ . One method of measuring distortions in the wavefront **130** is by determining a spatial separation  $\Delta z$  between a reference plane **131** (by way of example, a plane analogous to the ideal wavefront **110**) at a known distance  $Z_0$  from the eye **120** at each (x,y) point of the distorted wavefront **130** as the leading edge of the wavefront **130** traverses the distance  $z_0$ . This is described mathematically as:

$$\Delta z(x,y) = z_0 - W(w,y) \quad (1)$$

These  $\Delta z$  measurements define optical path differences due to aberrations in the eye **120** being tested, by way of example. An appropriate correction consists of removing these optical path differences. By way of example, such correction is performed at reference plane **131**.

Depending on the desired corrective therapy (corneal tissue ablation, synthetic lens addition, by way of example), the amount of material removed or added at each (x, y) coordinate can be calculated directly if the refractive index of the material in question is known. For many procedures, such as intra-ocular lens implantation or radial keratotomy, a wavefront analysis may be performed repetitively during a procedure to provide feedback information as to the appropriate endpoint of the procedure.

In terms of the illustrative example, the differences  $\Delta z(x,y)$  between the distorted wavefront **130** and the ideal wavefront **110** are the consequence of the aberrations in the eye. Correction of these aberrations consists of introducing an optical path difference at the reference plane **131** of negative  $\Delta z(x,y)$ . If the treatment approach, by way of example, consists of removing tissue from the surface of the cornea **126** by laser ablation, then one choice for the location of reference plane **131** is tangential to the surface of cornea **126** (i.e. at  $z=0$ ). This is illustrated schematically with reference to FIG. 1D, where the curvature of the cornea **126** is greatly exaggerated for clarity of illustration. Ablation is then carried out discretely at each (x,y) coordinate along the cornea **126** by a laser beam delivery and eye tracking system such as described in U.S. Patent Nos. 5,980,513; 5,849,006; and 5,632,742, commonly owned with the present invention, and which disclosure is herein incorporated by reference.

The appropriate corneal ablation depth at any (x,y) transverse coordinate is, to within a small error, given by:

$$\Delta z(x,y) / (n_c - 1) \quad (2)$$

where  $n_c$  is the refractive index of corneal tissue or 1.3775. The method described in detail below calculates  $\Delta z(x,y)$  by first measuring the local slopes in wavefront **130**, i.e.  $\partial W(x,y)/\partial x$  and  $\partial W(x,y)/\partial y$ , at a number of points in the transverse x and y directions in reference plane **131** and then generating a mathematical description of  $W(x,y)$  having slopes in best possible agreement with the experimentally determined values. One such slope  $\partial W(x,y)/\partial x$  is illustrated with reference again to FIG. 1D. In doing this, a small error is introduced due to the fact that distorted wavefront **130** is measured at the reference plane **131** while wavefront **130** emerged from a curved corneal surface just

posterior to reference plane **131**. By way of example, an error  $E_x(x,y)$  is the lateral displacement in the x-direction at each (x,y) location at the measurement plane (i.e., reference plane **131**) to the curved corneal surface. A similar error will be manifest for any corrections involving curved optical surfaces. The error will generally increase with both (x,y) displacement from the point of tangency and local wavefront error.

For refractive surgery, the error may be negligibly small. The magnitude of error  $E_x(x,y)$  can be found for each measurement location (x,y) measured at an arbitrary coordinate, e.g.,  $(x_0, y_0)$  by projecting that location back to the point of origin on the cornea **126**. This is explained mathematically with reference again to FIG. 1D, where by way of example, it is assumed that the error is in the plane of the figure, i.e., the plane defined by  $y=y_0$ , although it is quite straightforward mathematically to extend the analysis to include errors in the y-dimension. The quantification of a line L tracing the propagation of the wavefront **131** measured at  $(x_0, y_0)$  in the  $z_0$  reference plane from the corneal surface to the reference plane is:

$$L(x) = z_0 - \frac{(x - x_0)}{\partial W(x_0, y_0) / \partial x} \quad (3)$$

If the corneal surface in the plane of the figure is described by the expression  $S(x_0, y_0)$ , then the point of origin for the wavefront **131** in question can be found by finding the point of intersection between  $L(x)$  and  $S(x, y_0)$ . Mathematically, one finds the value  $x'$ , that satisfies  $L(x') = S(x_0, y_0)$ . The error  $E_x(x_0, y_0)$  is then given as  $E_x(x_0, y_0) = x' - x_0$ . Extending the analysis to consider errors in the y-direction would yield a similar expression for  $E_y$  where  $E_y(x_0, y_0) = y' - y_0$ . If significant, these transverse errors can be compensated for by laterally displacing the aberration correction calculated at each (x,y) coordinate by the amounts  $E_x(x,y)$  and  $E_y(x,y)$ .

In the case of human corneas, the transverse error under most circumstances will be negligible. The error will be zero at the origin where the corneal tissue and reference plane **131** are tangent. For human corneas, the tissue is approximately spherical with a radius of curvature of approximately 7.5-8.0 mm. The corrective treatment radius is typically no more than 3 mm, and local wavefront radius of curvature

will almost never exceed 50 mm (a 20 diopter refractive error). The transverse error E at a 3 mm treatment radius for a local wavefront radius of curvature of 50 mm is less than 40  $\mu$ m.

For certain ophthalmic procedures, wavefront analysis can also be used repetitively during the corrective procedure to provide useful feedback information. One example of such use would be in cataract surgery where wavefront analysis could be performed on the eye following placement of an intra-ocular lens implant (IOL). The analysis helps to identify whether the appropriate refractive power IOL has been inserted, or whether a different refractive power IOL should be used. Another example of repetitive wavefront analysis would be during keratoplastic procedures where the cornea of the eye is deliberately distorted by altering the mechanical tension around the periphery thereof. Here, repetitive wavefront analysis will be used to refine the degree of induced tension change at each point around the cornea thereby providing the tool to obtain optimum surface curvature for best visual acuity.

In order to perform wavefront analysis in a manner compatible with corrective procedures such as those described above, the amount of spatial separation of component portions of wavefront 130 relative to the corresponding component portions of the planar or ideal wavefront 110 is measured. It is the system and method of the present invention that allows such separation to be objectively and accurately measured for even substantially aberrated eyes 120 including those exhibiting severe defects such as severe myopia or hyperopia.

For the evaluation or measurement portion of the present invention, the patient's pupil should ideally be dilated to approximately 6 mm or more, i.e., the typical size of a human pupil in low light. Smaller amounts of dilation or no dilation at all may also be evaluated or measured. In this way, the eye is evaluated while it is using the greatest area of the cornea so that any correction developed from such measurement takes into account the largest usable corneal area of the patient's eye. A lesser amount of the cornea is used in daylight where the pupil is considerably smaller, e.g., on the order of 3 millimeters. Dilation can be brought about naturally by implementing the

measurement portion of the present invention in a low light environment such as a dimly lit room. Dilation can also be induced through the use of pharmacologic agents.

Referring now to FIG. 2, a simplified schematic of one exemplary embodiment of the apparatus **10** of the present invention is illustrated. The apparatus **10** includes a laser **12** for generating optical radiation used to produce a small-diameter laser beam **14**. The laser **12** generates a collimated laser light beam (represented by dashed lines for the beam **14**) of a wavelength and power that is eye-safe. For ophthalmic applications, appropriate wavelengths would include the entire visible spectrum and the near infrared spectrum. By way of example, appropriate wavelengths may be in a range of from approximately 400 to 1000 nanometers, including 550, 650, 850 useful wavelengths. While operation in the visible spectrum is generally desired, since these are the conditions in which the eye operates, the near infrared spectrum may offer advantages in certain applications. For example, the patient's eye may be more relaxed if the patient does not know measurement is taking place. Regardless of the wavelength of the optical radiation, power should be restricted in ophthalmic applications to eye safe levels. For laser radiation, appropriate eye-safe exposure levels can be found in the U.S. Federal Performance Standard for Laser Products. If the analysis is to be performed on an optical system other than the eye, the examination wavelength range logically should incorporate the intended performance range of the system.

To select a small-diameter collimated core of laser light beam **14**, an iris diaphragm **16** is used to block all of laser light beam **14** except for the laser beam **18** of a size desired for use. In terms of the present invention, the laser beam **18** will have a diameter in the range of approximately 0.5-4.5 millimeters with 1-3 millimeters being typical, by way of example. A badly aberrated eye uses a smaller diameter beam while an eye with only slight aberrations can be evaluated with a larger diameter beam. Depending on the output divergence of the laser **12**, a lens, as will be later described, can be positioned in the beam path to optimize collimating of the beam.

Laser beam **18**, as herein described by way of example, is a polarized beam that is passed through a polarization sensitive beam splitter **20** for routing to a focusing

optical train **22** which optical train operates to focus the laser beam **18** through the optics of the eye **120** (e.g., the cornea **126**, pupil **125** and the lens **124**) to the retina **122**. It is to be understood that the lens **124** may not be present for a patient that has undergone a cataract procedure. However, this does not affect the present invention.

5 In the illustrated example of FIG. 2, the optical train **22** images the laser beam **18** as a small spot of light at or near the eye's fovea centralis **123** where the eye's vision is most acute. Note that the small spot of light could be reflected off another portion of retina **122** in order to determine aberrations related to another aspect of one's vision. For example, if the spot of light were reflected off the area of the retina **122** surrounding the  
10 fovea centralis **123**, aberrations specifically related to one's peripheral vision could then be evaluated. In all cases, the spot of light may be sized to form a near-diffraction limited image on the retina **122**. Thus, the spot of light produced by laser beam **18** at fovea centralis **123** does not exceed approximately 100 micrometers in diameter and, typically, is on the order of 10 micrometers.

15 The diffuse reflection of the laser beam **18** back from the retina **122** is represented in FIG. 2 by solid lines **24** indicative of radiation that passes back through the eye **120**. The wavefront **24**, earlier described with reference to FIG. 1B as distorted wavefront **130** impinges on and is passed through the optical train **22** and on to the polarization sensitive beam splitter **20**. The wavefront **24** is depolarized relative to the  
20 laser beam **18** due to reflection and refraction as the wavefront **24** emanates from the retina **122**. Accordingly, the wavefront **24** is turned at the polarization sensitive beam splitter **20** and directed to a wavefront analyzer **26** such as a Hartmann-Shack (H-S) wavefront analyzer. In general, the wavefront analyzer **26** measures the slopes of wavefront **24**, i.e., the partial derivatives with respect to x and y, at a number of (x,y)  
25 transverse coordinates, as earlier described with reference to FIGS. 1C and 1D. This partial derivative information is then used to reconstruct or approximate the original wavefront with a mathematical expression such as a weighted series of Zernike polynomials.



The polarization states for the incident laser beam **18** and the beam splitter **20** minimizes the amount of stray laser radiation reaching the sensor portion of the wavefront analyzer **26**. In some situations, stray radiation may be sufficiently small when compared to the radiation returning from the desired target (e.g., the retina **122**) so that the polarization specifications are unnecessary.

The present invention is able to adapt to a wide range of vision defects and as such achieves a new level of dynamic range in terms of measuring ocular aberrations. Dynamic range enhancement is accomplished with the optical train **22** and/or a wavefront sensor portion of the wavefront analyzer **26**. With continued reference to FIG. 2, the optical train **22** includes a first lens **220**, a flat mirror **221**, a Porro mirror **222** and a second lens **224** all of which lie along the path of laser beam **18** and the wavefront **24**. The first lens **220** and the second lens **224** are identical lenses maintained in fixed positions. The Porro mirror **222** is capable of linear movement as indicated by arrow **223** to change the optical path length between the lenses **220** and **224**. However, it is to be understood that the present invention is not limited to the particular arrangement of the flat mirror **221** and the Porro mirror **222** and that other optical arrangements, as will herein be described by way of example, will be used without departing from the teachings and benefits of the present invention.

A "zero position" of the Porro mirror **222** is identified by replacing the eye **120** illustrated with reference again to FIG. 2, by a calibration source, as will be described later by way of further example, of collimated light to provide a reference wavefront such as the perfect plane wave **110**, earlier described with reference to FIG. 1A. Such a source could be realized by a laser beam expanded by a beam telescope to the diameter that will cover the imaging plane of wavefront analyzer **26** and adjustment of the Porro mirror **222** until the wavefront analyzer **26** detects the light as being collimated. Note that the changes in optical path length brought about by the Porro mirror **222** can be calibrated in diopters to provide an approximate spherical dioptric correction, as will be explained further below.

The dynamic range of the apparatus **10** is further improved by providing an improved wavefront sensor arrangement **28** as illustrated with reference to FIGS. 3 and 4. The wavefront analyzer **26** includes an opaque imaging plate **32** having an array of holes **34** passing therethrough, a planar array **36** of light-sensitive cells such as charge coupled device (CCD) cells **38**, and a processor **40** operable with the planar array **36** of the CCD cells **38**. The combination of the plate **32** and the planar array **36** provides one embodiment of the present invention. The plate **32** is held parallel to and spaced from the planar array **36** by a separation distance  $F$ . As will be explained further below, the separation distance  $F$  can be varied to adjust for signal gain. To do this, the planar array **36** is coupled to a positioning apparatus **42**, e.g., a conventional motorized linear positioner having precise movement capability, that adjusts the position of planar array **36** relative to the plate **32** for changing the separation distance  $F$  as indicated by arrow **43**. With respect to the array of holes **34**, each of the holes **34** are of equal size and shape with a circle being typical owing to its ease of manufacture. As herein described by way of example, a square array geometry is used for the array of holes **34**, although other array geometries will be used without departing from the teachings of the present invention.

As illustrated with reference to FIG. 4, when the wavefront **24** impinges on the plate **32**, a portion of the wavefront **24**, indicated by arrow **25**, passes through the hole **34** to illuminate planar array **36**. To a first order, the resulting image formed by each such wavefront portion **25** is a positive shadow of the respective hole **34**. However, diffraction occurs as determined by the diameter  $D$  of each hole **34**, the wavelength  $\lambda$  of the light source (e.g. the wavefront **24**) and the separation distance  $F$  between the plate **32** and the planar array **36**. The value of  $F$  is varied by the positioning apparatus **42** to adjust the gain based on the particular patient as will be explained further below.

Note that performance of the plate **32** with holes **34** may also be accomplished using a solid plate or film made from a light-sensitive material such as a photo-lithographic film. In such a case, the array of holes **34** would be replaced by an array of shaped light transmissive apertures through which light passes when impinging

thereon. The remainder of such a plate or film would be impervious to light. Such an embodiment permits the light transmissive apertures to be easily made to conform to any desired shape.

Regardless of how each wavefront portion **25** is generated, the present invention  
5 measures the amount of angular deflection of each wavefront portion **25** relative to a  
wavefront portion **112** that results from a calibration wavefront such as the planar  
wavefront earlier described. The calibration or planar wavefront of light results in the  
wavefront portion **112** impinging at a normal or perpendicular to plate **32** and  
10 illuminates a geometric spot **114** on the planar array **36**. In contrast, continuing with the  
wavefront **24** representing a distorted wavefront as described above, the wavefront  
portion **25** will exhibit an amount of angular deflection relative to the calibration  
wavefront portion **112**. The angular deflection causes the wavefront portion **25** to  
illuminate a geometric spot **27** on the planar array **36** that is offset from the spot **114**. In  
15 terms of the present invention, the amount of offset is measured relative to centroids  
**116** and **29** of spots **114** and **27**, respectively. In the two dimensions of the planar array  
**36**, the centroid **29** is typically deflected in both the x and y directions of the array **36**.  
Thus, the angular deflection in each of the x and y directions is given by  $\Delta x/F$  and  $\Delta y/F$ ,  
respectively.

With reference again to FIG. 2, the lenses **220** and **224** in one embodiment are  
20 identical as mentioned above. However, in certain applications it may be desirable to  
magnify or minify the wavefront at the wavefront sensor. This can be accomplished by  
using lenses **220** and **224** of different focal lengths and adjusting dimensions of the  
apparatus **10** accordingly. For ophthalmic evaluation, the object plane of the apparatus  
should ideally be tangent to the corneal surface which can be achieved by a variety of  
25 means. Thus, each point at the object plane of the optical train **22** very nearly  
corresponds to the same point on the cornea **126**. However, since the cornea **126** is  
curved, there will be a slight lateral displacement. The plate **32** described earlier with  
reference to FIG. 4 of the wavefront analyzer **26**, or an imaging plane of any wavefront  
sensor portion, is positioned at the focal plane of lens **220**. In this way, the object plane

is always imaged on the plate **32** in direct correspondence with the wavefront image emerging from cornea **126**. This will be true regardless of the optical path length between the lenses **220** and **224**. There are several advantages to this structure, one of which is that there are very good planar arrays of light-sensitive cells that are commercially available to image an area corresponding to the 6 millimeter central circular region of the cornea.

The plate **32** (or the imaging plane of any wavefront sensor portion of a wavefront analyzer) breaks the wavefront **24** into wavefront pieces that can each be measured independently in terms of propagation direction at the planar array **36**. Since in an embodiment herein described by way of example, the optical train **22** does not magnify or reduce the image in the object plane, a point at the object plane corresponds to the same point at the image plane of the optical train. With the Porro mirror **222** set at its zero position, the direction each portion of the wavefront **24** is traveling toward the object plane is reproduced exactly at the image plane of wavefront analyzer **26**. By way of example, if one wavefront portion at a location in the object plane was traveling away from the optical axis at an angle of  $20^\circ$  with respect to the optical axis that is perpendicular to the object plane, the wavefront portion at the same location in the image plane will also be traveling away from the optical axis at an angle of  $20^\circ$ .

Note that a person who is myopic will produce a wavefront such that the wavefront portions/pieces isolated by the plate **32** will converge toward the center of planar array **36**. A hyperopic person will produce a wavefront such that the wavefront pieces isolated by the plate **32** diverge. Thus, a person with a significant vision error becomes difficult to evaluate because wavefront portions can either overlap (myopia) at the planar array **36** or spill off (hyperopia) the planar array.

In the present invention, five ways of compensating for such severe aberrations are herein described by way of example. The first way is to utilize a wavefront sensor with sufficiently small light sensitive cells **38** and sufficiently large holes **34** (or any other transmissive aperture). In this way, measurement of each wavefront piece can be performed to an acceptable accuracy using a small value for  $F$ . A second way is to

move planar array **36** along the optical axis to change the separation distance  $F$  to the plate **32**. For a person with a severe aberration, the planar array **36** is positioned close to the plate **32** to keep the projected wavefront portions well separated and on the planar array. For a mild aberration, the planar array **36** is moved to increase the separation distance  $F$  to the plate **32** to make a more accurate measurement. The advantage of moving the planar array **36** to change the separation distance  $F$  to the plate **32** is that the wavefront analysis is easily achieved for any position. Yet another way of compensating for severe aberrations using the present invention is to change the optical path length between lenses **220** and **224**. Moving the Porro mirror **222** will not affect where the wavefront hits the plate **32**, but will change the angular deflections at which the projected wavefront portions pass through the plate **32**, i.e.,  $\Delta x/F$  and  $\Delta y/F$ . Decreasing the optical path length between lenses **220** and **224** will tend to pull the wavefront portions toward the center of planar array **36** thereby compensating for hyperopia. Increasing the optical path length between lenses **220** and **224** will tend to spread the wavefront portions toward the edges of the planar array **36** thereby compensating for myopia. The degree to which the angular deflection associated with each wavefront piece is altered is a linear function of its distance off the optical axis and the movement of the Porro mirror **222** from its zero position. A fourth way of compensating for severe aberrations is to insert one or more trial lenses of specified sphero-cylindrical power at the location of the intermediate focal plane, as will be discussed in detail later in this section. This serves to reduce or remove low order aberrations from the wavefront so that displacement of spots at the CCD cells **38** is minimized and accurate evaluation can proceed. The effect of the specified lens addition is then included in the final wavefront reconstruction. A fifth way is to increase the magnification of the wavefront at the wavefront sensor relative to that at the eye. This is accomplished by an appropriate choice of lenses in the relay optic design. Magnification will reduce the slope of the wavefront uniformly, thereby reducing the displacement of each spot at the CCD.

By way of example, to accurately determine the centroid **29** of the spot **27** of light impinging on the planar array **36**, a fine structure of cells **38** relative to a spot size is provided. Each spot covers a plurality of cells **38**. One method used to determine the centroid **29** of each spot **27** unambiguously with respect to a spot caused by another one of the holes **34**, assigns a unique number of cells **38** to each hole **34**. The "assigned areas" are designated, as illustrated with reference to FIG. 5, by way of example, with the heavy grid lines **39**. It is to be understood that the grid lines **39** are not actual physical boundaries between cells **38** but are shown simply to illustrate the unique designated areas containing a plurality of the cells **38**. It is anticipated that other centroid strategies will be utilized that do not necessitate such partitioning of the array **36** given the teachings of the present invention. An alternative method for identifying and correlating centroids is later described in this section.

By way of example, the present invention could also be practiced with a wavefront analyzer that replaces plate **32** described with reference to FIG. 3, with a two dimensional array of identical spherical lenslets **33**, as illustrated with reference to FIG. 6. In such an embodiment, the lenslet array **33** may be operable by the positioning apparatus **42** such that separation distance  $F$  is independent of the focal length  $f$  that defines the focal plane of the lenslet array **33** which is represented by dashed line **35**. Each wavefront portion **37** passed through a sub-aperture of the lenslet array **33** is reduced in size (e.g., diameter) but is not necessarily brought to a minimum focus at the planar array **36** as it would be if separation distance  $F$  were equal to focal length  $f$ . In the operation of this embodiment configuration, the lenslet array **33** is positioned to concentrate the light in each wavefront portion of an area for providing sufficient intensity onto the planar array **36**, yet still illuminating a substantial plurality of cells **38** for greatest accuracy in determining the deflection of the centroids **29**.

Regardless of the structure of the wavefront sensor, the processor **40** computes each two-dimensional centroid **29** of each spot **27** generated by the wavefront **24**. The amount of two dimensional centroid shift relative to the centroid of the calibrating spot for each designated area associated with a corresponding hole **34** (or sub-aperture of

lenslet array **33**) is divided by the separation distance  $F$  to generate a matrix of local slopes of the wavefront, i.e.,  $\partial W(x,y)/\partial x$  and  $\partial W(x,y)/\partial y$  at the  $(x,y)$  coordinates of the centers of holes **34**. For simplicity of discussion, these will be indicated by  $P(x,y)=\partial W(x,y)/\partial x$  and  $Q(x,y)=\partial W(x,y)/\partial y$ , respectively.

Numerous methods exist for using the partial derivative data to calculate the distorted wavefront **130** and **24** as described above with reference to FIGS. 1B and 2. One acceptable approach is that described by Liang et al. in the aforementioned Journal of the Optical Society of America paper, where the wavefront is approximated using Zernike polynomials. This is a standard analytic technique described in numerous optics texts such as "Principles of Optics, 11 by M. Born and E. Wolf, Pergamon Press, Oxford, England, 1964. By way of example, the Zernike polynomial approach will be discussed herein. However, it is to be understood that other mathematical approaches can be used in approximating the distorted wavefront. It will be understood by one of ordinary skill in the art that other mathematical approaches can be used in approximating the distorted wavefront. By way of example, such approaches may include the use of Fourier series and Taylor series.

$$W(x,y) = \sum_{i=0}^n C_i Z_i(x,y) \quad (4)$$

Briefly, the wavefront  $W(x,y)$  is expressed as a weighted sum of the individual polynomials where  $C_i$  are the weighting coefficients, and  $Z_i(x,y)$  are the Zernike polynomials up to some order. The upper limit  $n$  of the summation is a function of the number of Zernike polynomials, i.e., the highest order, used to approximate the true wavefront. If  $m$  is the highest order used, then

$$n = (m+1)(m+2)/2 \quad (5)$$

Derivation of the Zernike polynomials up to an arbitrary order  $n$  is described in numerous optical texts such as the aforementioned book by Born and Wolf. One possible method of determining the centroid **29**, **116** of a spot **27**, **114**, respectively, as earlier described with reference to FIGS. 4 and 5, and calculation of the Zernike weighting coefficients will now be explained. The directions of the unit normals at the center of each hole **34** are based on the centroids of the spots on cells **38**.

Since each spot will illuminate a plurality of cells varying intensity, a standard amplitude-weighted centroid calculation can be used to find the center of each spot. In order to clearly delineate each centroid from background noise, by way of example, resulting from spurious light reaching the CCD surface between valid spots, standard mathematical techniques such as a matched spatial filter can be applied to the CCD data prior to centroid identification.

An alternative method is herein described for identifying individual spots and correlating their geometry. The apparatus is configured such that the optical axis is aligned to the center of a particular aperture at the entrance face of the wavefront sensor. This aperture is located at or near the center of the entrance face. If the probe beam entering the eye is also aligned to the system optical axis, then due to the reversible nature of light rays, a light spot will always be seen directly behind the aligned aperture. That is, a spot will always be seen on the CCD sensor at this location, regardless of the wavefront aberrations, and will always correspond to the overlying aperture. Immediately adjacent spots will be minimally displaced from their "zero slope" locations. As one moves further from the central reference spot, generally greater spot displacements will occur. Using this knowledge, it is a relatively straight forward process to identify all the spots in the CCD pattern and establish their geometric relationships.

The displacement of the centroid from that of a perfectly collimated light beam, corresponding to ideal and emmetropic vision, is then calculated and used to determine the wavefront slope at each sample location. The location of the centroids for a collimated light beam may either be directly measured in a calibration step prior to the patient exam, or taken from a calculated reference pattern based on the wavefront sensor construction.

Multiple exposures may be used to check for improper eye alignment or eye movement during individual exposures. If eye movement during exposures cannot be analyzed successfully by acquiring multiple exposures, then the apparatus 10 can be augmented by the addition of an eye tracker 30, illustrated with reference again to FIG. 2. One possible placement of the eye tracker 30 is herein illustrated. However, it is to



be understood that the eye tracker **30** could be placed elsewhere within the apparatus **10**. One such eye tracker is disclosed in the aforementioned U.S. Patent No. 5,980,513, commonly owned with the present invention. In this way, wavefront analysis is performed even during a limited amount of eye motion.

A one-time calibration exposure can also be used to determine the relative sensitivities of the individual cells. This is made in uniform collimated light with plate **32** removed. The responses of individual cells are then recorded. For each light transmissive aperture (e.g, hole **34**), the centroid in the collimated case serves as a dedicated origin for the particular hole. The shift from the "origin" for each hole to the centroid caused by the wavefront **24** (as observed in this coordinate system) is determined by the direction of the wave surface corresponding to that hole. If  $\Delta x(m,n)$  is the x-component of the (m,n)th centroid and F is the plate separation, then the P-value for the (m,n)th centroid is:

$$P(m,n) = \partial x(m,n) / \partial z = \Delta x(m,n) / F \quad (6)$$

The corresponding expression for Q is:

$$Q(m,n) = \partial y(m,n) / \partial z = \Delta y(m,n) / F \quad (7)$$

Thus, each P(m,n) and Q(m,n) represents the partial derivatives of W(x,y) with respect to x and y for the (x,y) coordinates of each hole **34**. For an m-order Zernike approximation of the original wavefront, the experimentally determined P's and Q's are then used in the following equations to calculate the appropriate  $C_i$  weighting coefficients as follows:

$$P(m,n) = \frac{\partial W(x,y)}{\partial x} = \sum_{i=0}^n C_i \frac{\partial Z_i(x,y)}{\partial x} \quad (8)$$

$$Q(m,n) = \frac{\partial W(x,y)}{\partial y} = \sum_{i=0}^n C_i \frac{\partial Z_i(x,y)}{\partial y} \quad (9)$$

By using a least-squares approx(m,n)/∂z each to minimize the error between the actual wavefront slopes on the left hand side in the above equations and the Zernike approximations on the right hand side, optimal values for the weighting coefficients can be obtained.

In one possible approach to calculating a centroid  $(x_c, y_c)$ , each hole **34** is assigned its dedicated area of the array **36** or  $(i_{m,n} \pm \Delta i, j_{m,n} \pm \Delta j)$ . This square of many light-sensitive cells is large enough that neighboring hole images never encroach, and all illumination from this hole is contained. The square contains  $4\Delta i \cdot \Delta j$  cells.

If array 36 is designated  $C_{k,1} = (x_c(i, j), y_c(i, j))$ ,  $k, 1 = 0 \dots 2\Delta i, 2\Delta j$ , and the spacing on centers is  $\Delta x = \Delta y = d$ , the measured cell responses are  $V(k, 1)$  and the relative responsivities are  $R(k, 1)$ , then the x-component  $x_c$ , a function of  $i, j$  is represented by

$$x_c(i, j) = \left[ \sum_{k,1} V(k,1) * R(k,1) * d * k \right] / \left[ \sum_{k,1} V(k,1) * R(k,1) \right] \quad (10)$$

and the y-component  $y_c$ , as a function of  $i, j$  is represented by

$$y_c(i, j) = \left[ \sum_{k,1} V(k,1) * R(k,1) * d * l \right] / \left[ \sum_{k,1} V(k,1) * R(k,1) \right] \quad (11)$$

Then, if  $(x_{c0}(i, j), y_{c0}(i, j))$  is the "origin centroid" for the  $(i, j)$  hole, i.e., made in perpendicular collimated light, and  $(x_{cw}(i, j), y_{cw}(i, j))$  is the corresponding centroid found for the wavefront to be measured, then the relative centroid shift  $(x_{cr}(i, j), y_{cr}(i, j))$  is found as

$$x_{cr}(i, j) = x_{cw}(i, j) - x_{c0}(i, j) \quad (12)$$

$$y_{cr}(i, j) = y_{cw}(i, j) - y_{c0}(i, j) \quad (13)$$

The values  $P(i, j)$  and  $Q(i, j)$  are determined from

$$P(i, j) = x_{cr}(i, j) / F \quad (14)$$

and

$$Q(i, j) = y_{cr}(i, j) / F \quad (15)$$

The surface partial derivatives  $P(i, j)$  and  $Q(i, j)$  for the array of hole centers of plate 32 are next used to calculate the appropriate Zernike polynomial weighting coefficients to describe the original wavefront  $W(x, y)$ . This will now be explained by way of illustration for a 7 x 7 square array of holes 34. However, it is to be understood that other sizes and shapes of hole arrays could be used.

First, a 1 x 98 matrix (i.e., column vector)  $PQ(k)$  is formed as

$$PQ(k) = P(7i + j), j = 0...6, i = 0...6, k = 0...48 \quad (16)$$

$$PQ(k) = Q(7i + j), j = 0...6, i = 0...6, k = 49...98 \quad (17)$$

with  $j$  cycling for each  $i$ , i.e.,  $PQ(18) = P(2, 5)$ .

The matrix  $PQ$  is multiplied from the left with a transition matrix  $TM$  to get the matrix  $C$  as follows

$$C = TM * PQ \quad (18)$$

where  $TM$  is a 98 wide by 14 high matrix and  $C$  is a 1 wide by 14 high matrix or column vector.  $C$  is the matrix  $C_k$   $k=1, \dots, 14$  such that, to a least square error,

$$W(x, y) = \sum_k C_k * Z_k(x, y) \quad (19)$$

and  $TM$  is calculated for a given aperture, e.g., a 6 millimeter pupil aperture. The functions  $Z_k(x, y)$  in equation (19) are the Zernike polynomials. There is no standard convention as to their sequence. Thus, for consistency, it is important that the same sequence is used to produce the set  $C_k$  that was chosen for deriving the matrix  $TM$ . They occur in groups of the same order, which is the highest exponent in the group, with the total number of members in an order increasing with the order. For example, in a fourth order analysis, orders up to and including 4 are used (less  $Z_0$  - the single member of order 0 that is the constant 1 which describes the reference position of the group in the  $z$  direction). Since wavefront 24 is moving along  $z$  (at the velocity of light),

this "piston term" describes only an arbitrary offset in Z, and this term may be ignored. The first 5 orders (0, 1, ...,4) contain 15 functions including the piston term.

Thus, in the illustrated example, 14 values of  $C_k$  are calculated as coefficients of 14 Zernike polynomials. By way of example, one such order used to calculate TM is herein illustrated, and includes both the Zernike functions and their partial derivatives.

## ZERNIKE (X,Y) POLYNOMIAL EXPANSION THROUGH ORDER 4

### Polynomial Order 0

	Z(0)	+1
10	dZ(0)/dx	0.0
	DZ(0)/dy	0.0

### Polynomial Order 1

	Z(1)	+y
	dZ(1)/dx	0.0
15	dZ(1)/dy	+1
	Z(2)	+x
	dZ(2)/dx	+1
	dZ(2)/dy	0.0

### Polynomial Order 2

20	Z(3)	-1+2y <sup>2</sup> +2x <sup>2</sup>
	dZ(3)/dx	+4x
	dZ(3)/dy	+4y
	Z(4)	+2xy
	dZ(4) /dx	+2y
25	dZ(4) /dy	+2x
	Z(5)	-y <sup>2</sup> +x <sup>2</sup>
	dZ(5) /dx	+2x
	dZ(5) /dy	-2y

### Polynomial Order 3

30	Z(6)	-2y+3y <sup>3</sup> +3x <sup>2</sup> y
----	------	--

	$dZ(6)/dx$	$+6xy$
	$dZ(6)/dy$	$-2+9y^2+3x^2$
	$Z(7)$	$-2x+3xy^2+3x^3$
	$dZ(7)/dx$	$-2+3y^2+9x^2$
5	$dZ(7)/dy$	$+6xy$
	$Z(8)$	$-y^3+3x^2y$
	$dZ(8)/dx$	$+6xy$
	$dZ(8)/dy$	$-3y^2+3x^2$
	$Z(9)$	$-3xy^2+x^3$
10	$dZ(9)/dx$	$-3y^2+3x^2$
	$dZ(9)/dy$	$-6xy$
	<u>Polynomial Order 4</u>	
	$Z(10)$	$+1-6y^2+6y^4-6x^2+12x^2y^2+6x^4$
	$dZ(10)/dx$	$-12x+24xy^2+24x^3$
15	$dZ(10)/dy$	$-12y+24y^3+24x^2y$
	$Z(11)$	$-6xy+8xy^3+8x^3y$
	$dZ(11)/dx$	$-6y+8y^3+24x^2y$
	$dZ(11)/dy$	$-6x+24xy^2+8x^3$
	$Z(12)$	$+3y^2-4y^4-3x^2+4x^4$
20	$dZ(12)/dx$	$-6x+16x^3$
	$dZ(12)/dy$	$+6y-16y^3$
	$Z(13)$	$-4xy^3+4x^3y$
	$dZ(13)/dx$	$-4y^3+12x^2y$
	$dZ(13)/dy$	$-12xy^2+4x^3$
25	$Z(14)$	$+y^4-6x^2y^2+x^4$
	$dZ(14)/dx$	$-12xy^2+4x^3$
	$dZ(14)/dy$	$+4y^3-12x^2y$

30 The choice of sequencing the Zernike polynomials dictates the interpretations of the  $C_k$  in equation (19) and therefore the order of terms in the TM matrix. Hence, the

TM matrix is calculated after the choice is made. The development of the TM matrix for the illustrated example will be explained below.

Note that the fourth order analysis is only an example and is not the only possibility. A Zernike analysis can be done to any order. In general, the higher the order, the more accurate the result over the tested points. However, an exact polynomial fit over the tested points is not necessarily desirable. Such fits have the typical disturbing property that, unless the surface itself happens to be an exact polynomial of order no higher than that used for the surface fit, forcing an exact fit at separated points often causes wild swings between fitted points. That is, in polynomial surface fitting, an exact fit at a finite number of points can yield a poor average fit for a general function.

Calculation of the  $\Delta z(x,y)$  optical path difference information from the Zernike reconstruction of the wavefront is accomplished simply by subtracting a constant from the Zernike approximation. The value of the constant will depend on the desired characteristics of  $\Delta z(x,y)$ . Depending on the method chosen to correct the aberrations (e.g., laser ablation, lens addition, etc.) it may, for example, be desirable to set either the maximum, mean or minimum value in  $\Delta z(x,y)$  equal to zero.

The development of the transition matrix TM will now be explained for the illustrated example of a 7 x 7 array of holes in plate 32. At each point  $(x_i, y_j)$ , the tangents of the components of the normal are  $P(x_i, y_j)$  and  $Q(x_i, y_j)$  where

$$P(x_i, y_j) = \partial W(x_i, y_j) / \partial x \quad (20)$$

and

$$Q(x_i, y_j) = \partial W(x_i, y_j) / \partial y \quad (21)$$

Combining these with equation (11),

$$P(x_i, y_j) = \sum_k C_k \partial W(x_i, y_j) / \partial x \quad (22)$$

and

$$Q(x_i, y_j) = \sum_k C_k \partial W(x_i, y_j) / \partial y \quad (23)$$

each applicable to 49 (i,j) combinations. These are combined into a single column vector PQ that is 98 elements high, i.e., a 98 x 1 matrix. Defining two matrices  $C_k$  (14 high x 1 wide) and  $M_{k,(i,j)}$  (14 wide x 98 high)

$$\left( M_{k,(i,j)} \right) = \partial Z_k(x_i, y_j) / \partial x \quad ; \quad \partial Z_k(x_i, y_j) / \partial y \quad (24)$$

where the x-derivatives are the first 49 rows and the y-derivatives are the last 49 rows.

5 Then, equation (19) can be rewritten as the matrix equation

$$(PQ) = (M)(C) \quad (25)$$

where the top 49 rows of M are the  $\partial W(x_i, y_j) / \partial y$ .

The expression in equation (25) gives the normal components in terms of the Zernike coefficients for a surface described by the array of 14 C's. These are exact, but it is not guaranteed that the actual total surface can be described by such an array of coefficients. Accordingly, if it is assumed that the description is within an acceptable tolerance, i.e., tolerating the errors that remain after least square error determination, then equation (26) can be considered to define the column vector C implicitly in terms of the mathematical matrix M and the measured vector PQ, both of which are known. The method of effecting the solution under the minimization condition is as follows.

15 First, equation (25) is multiplied on the left by  $M^T$ , the transpose of M such that

$$(M^T)(PQ) = (M^T)(M)(C) = (S)(C) \quad (26)$$

where

$$S \equiv M^T M \quad (27)$$

is a square and symmetric matrix, e.g., of dimensions 14 x14 (with each element the sum of 98 products). Such a matrix has an inverse unless the determinant of its coefficients is zero. Since this is based on the Zernike polynomials alone, and they are

all independent of each other, the determinant is non-zero, so that an inverse  $S^{-1}$  is defined. Next, equation (25) is multiplied on the left by  $S^{-1}$  to yield

$$(S^{-1})(M^T)(PQ) = (S^{-1})(S)(C) = (I)(C) = C \quad (28)$$

Then, the mathematical transition matrix (independent of measurement) is

$$(TM) = (S^{-1})(M^T) \quad (29)$$

and the "best fit" array of C's from the measured PQ's can be produced by the simple matrix multiplication

$$(C) = (TM)(PQ) \quad (30)$$

To evaluate the eye unambiguously, all spots illuminating the planar array **36** due to a wavefront **24** are incident on the planar array simultaneously. If it is desired to reduce effects of eye movement, a pulsing or shuttering laser source may be used, or an eye tracker.

An implementation of the present invention suitable for clinical use is illustrated, by way of example, with reference to FIG. 7 and is referenced generally by numeral **11**. Like reference numerals are used to describe elements that are the same as those described above with respect to the apparatus **10**. A dichroic beam splitter **52** is interposed between the beam splitter **20** and the optical train **22** to introduce fixation target optics **60** and observation optics **70** into the apparatus **11** which are optically separated from one another by a 50/50 beam splitter **54**. Fixation target optics provide the eye **120** with visible light in the shape of a target. The visible light generated by fixation target optics **60** is reflected by the dichroic beam splitter **20** and directed through optical train **22**.

It is to be understood that the fixation target optics **60** can be implemented in a variety of fashions. By way of example, one such embodiment is shown and includes a visible light source **61**, a light diffuser **62**, a target **63**, a field stop **64**, a lens **65** and an iris **66**. The light source **61** and the light diffuser **62** are used to provide uniform illumination of the fixation target **63**. The field stop **64**, lens **65**, and iris **66** are used in



conjunction with the optical train **22** to present a desired image of the fixation target **63** to the patient for viewing by the eye **120**.

Observation optics **70** allows a technician to view and document an eye evaluation procedure. While a variety of implementations of observation optics **70** are possible, one such implementation is shown by way of example, with reference again to FIG. 7. The observation optics **70** includes a field lens **71**, lens **72**, iris **73**, lens **74**, and a camera **75**. A ring illuminator **80** is placed in front of the eye **120** to illuminate same for observation and/or filming purposes.

With reference now to FIGS. 9-11, an exemplary embodiment of the apparatus **10** will be herein described beginning with series **300**, which improved apparatus **300** is constructed as a patient examination station which allows the patient **302** to be comfortably positioned for the measurement of the eye **120**, as earlier described. For convenience to the technician operating the apparatus **300**, a computer monitor, mouse, and keyboard are located on a separate cart for this embodiment of the present invention, herein described. The apparatus **300** includes a housing **304** having a platform **306** which is carried by a rigid frame **308**. The frame **308** includes wheels **310** to facilitate shipping and installation at the clinical site, as well as locking and leveling feet **312** for securing the apparatus to the supporting floor **314**. Once the apparatus is positioned, the integrated leveling feet **312** are deployed to provide a stable stationary frame **308**, and thus platform **306**.

As illustrated, by way of example with reference again to FIGS. 9-11, the patient **302** sits at a patient end **316** of the apparatus **300**, with his or her head resting in a headrest **318**, which headrest is adjustable in directions left/right (X-direction), up/down (Y-direction), or toward/away (Z-direction) relative to the platform **306**, using adjustment assembly **320**. The headrest **318** is attached to the lower surface of the platform **306** which forms an optical table for mounting optical components thereon, as illustrated with reference to FIG. 12, and as will herein be described in further detail. The housing **304** includes a removable optical table cover **322** which protects the optical components carried within the housing. The optical table cover **322** is secured to the

platform **306** with keyed locks to prevent unauthorized access to the optical components. The platform **306** is bolted to the rigid frame at four locations **307**, as illustrated with reference again to FIGS. 9 and 10. The frame **308** also carries electronics **324** and a computer **326** which includes the processor **40** earlier described with reference to FIG. 6, as well as a connector plate for a computer keyboard, monitor and mouse. The frame **308** also includes an upper bay **328** housing electronics controlling optical components carried by the platform **306**, and a lower bay **330** housing an uninterruptible power supply (UPS) and an isolation transformer.

As illustrated with reference again to FIGS. 9-12, three ports are positioned within the cover **322**, and include an examination port **332** to allow the wavefront measurement of the eye **120** to take place, and two eye illumination ports **334** which allow lamps **336** carried within the housing **304** to illuminate the eye for visualization by an internal video camera **338**. In addition, the adjustment assembly **320** includes a position sensor **321** which senses an x-direction displacement for detecting a position of the headrest **318** to the left or to the right of a reference center line location **3191**. A signal indicative of the sensed displacement is provided to the computer **326** for automatically recording the appropriate eye **120** (e.g. left or right) being measured.

As illustrated with reference again to FIG. 12, the platform **306** provides an optical table with the patient positioning the eye **120** for measurement by the apparatus **300**. The platform surface measures approximately two feet by four feet, with the optical components fixed to the surface using a combination of commercial and customized precision hardware mounts. All transmissive optical elements have surface anti-reflection coating optimized for the selected probe beam wavelength. The optical layout includes five distinct optical pathways which share the optical elements as will herein be described, by way of example. With reference again to FIG. 12, a first optical path **340**, a fixation target image optical path illustrated in isolation in FIG. 12A for convenience to the reader, displays a fixation target image to the patient seated at the apparatus **300**. The patient aligns his/her visual axis to the optical axis **342** by looking at the center of a target reticle **344** having a grid pattern. With reference again to FIG.

12, a second optical path **346**, a video image optical path illustrated in isolation in FIG. 12B for convenience to the reader, captures a video image of the corneal plane. This allows the technician to assist in aligning the eye **120** for examination, and to record the exact location of the eye during each measurement using software reticles  
5 superimposed on a video image. With reference again to FIG. 12, a third optical path **348**, a probe laser optical path illustrated in isolation in FIG. 12C for convenience to the reader, sends a probe laser beam **350** into the eye **120** along the optical axis **342**. As earlier described with reference to FIGS. 2 and 7, the probe laser beam **14**, herein referred to with numeral **350** is attenuated to an eye-safe intensity and linearly polarized  
10 before being focused onto the corneal surface. With reference again to FIG. 12, a fourth optical path **352**, a re-emitted wavefront optical path illustrated in isolation in FIG. 12D for convenience to the reader, conveys the reflected wavefront **24** of FIG. 2, and herein described with numeral **354** re-emitted from the eye **120** and directed towards a wavefront sensor **356**. To accomplish this, first and second afocal relay stages **358**,  
15 **360** transfer the reflected wavefront **354** from the corneal plane of the eye **120** to the entrance face of the wavefront sensor **356**. Finally, with reference again to FIG. 12, a fifth optical path **362**, a calibration wavefront optical path illustrated in isolation in FIG. 12E for convenience to the reader, injects collimated laser light into the wavefront transfer path leading to the sensor **356**. Software operable within the computer **326**,  
20 described earlier with reference to FIG. 9, uses collimated light wavefront sensor output data to calibrate the apparatus **300** prior to patient measurement.

With continued reference to FIGS. 12, and 12A, the first optical path **340** is herein described as a fixation path which provides a reference image to the patient, such that the eye **120** is properly aligned when the patient is fixating on the reticle **344**  
25 of a reference target **366**. A target illumination lamp **368** back-lights the fixation target **366**, which fixation target image reaches the patient eye **120** by transmission through a 50/50 beam splitter **370**, lenses **372**, reflection in 50/50 beam splitters **374**, **376**, and transmission through lens combinations of afocal relay stage **358**, as well as through polarizing beam splitter **378**. In addition, a spectral filter is placed over the target

illumination lamp **368** to remove radiation over the 620-790 nm wavelength range that might otherwise interfere with a wavefront measurement at 670 nm. The lens combinations in the first relay stage **358** contain identical lens elements mounted in reverse order. Each consists of two meniscus lens elements, with an interposed achromatic doublet. The lens combinations work in tandem as a unity magnification afocal relay stage.

The optical elements including the polarizing beam splitter **378**, the lenses of the first afocal stage **358**, the beam splitters **374**, **376**, and one lens **380** of the lenses **372** are mechanically fixed in place on the surface of the platform **306**. The optical elements including a lens pair **382** of the lenses **372**, the beam splitter **370**, the fixation target **366**, and the illumination lamp **368** are all mounted on one precision linear translation stage, capable of movement along the optical axis **342** of this pathway. Translation of these optical elements focuses the fixation target **366** for the patient's view, compensating for any myopia/hyperopia present in the eye **120**. During patient examination the focus translation stage is adjusted to place the target optically just beyond the eye's infinity focal plane. This allows the patient to see a relatively distinct reticle pattern without stimulating accommodation by the eye **120**. The beam splitters **378**, **376**, **374** serve as interfaces between other optical pathways within the optical axis **342**, as will herein be described in further detail. By way of example, the beam splitter **370** is included for alignment purposes. A photo-detector **384** attached to the center of the left edge of beam splitter **370** senses light transmitted toward the fixation target along the optical axis.

With reference again to FIGS. 12 and 12B, the second optical path **346** captures video images of the eye **120** at an examination plane. This allows the clinical operator/technician to assist in patient alignment, and to measure actual eye displacement during the wavefront measurement. As earlier described, the illumination lamps **336** illuminate the eye **120**. The image of the eye is conveyed to the video camera **338** by transmission through the polarizing beam splitter **378** and the lens combinations **358**, reflection in the 50/50 beam splitter **376**, transmission through the

50/50 beam splitter **374**, reflection off mirror **386**, and transmission through lens **388**.

All these optical elements are fixed in place on the surface of the platform **306**. By way of example, this second path **346** provides a video field of view approximately 22 mm in diameter at the eye plane, with a limiting resolution of ~64 mm. As earlier described, a number of filters are placed in front of each eye illumination lamp **336** to reduce the spectral bandwidth of the radiation reaching the eye **120**. By way of example, these will include a blue filter to remove light at wavelengths below ~455nm (for eye safety), an infrared filter to remove light at wavelengths above ~920nm (for eye safety), and a rejection filter to remove light over the wavelength range 620nm - 790nm (to prevent interference with the wavefront measurement at 670 nm).

With continued reference to FIGS. 12 and 12C, the third optical path **348** irradiates a small spot on the patient's retina with eye safe laser radiation, as earlier described with reference to FIGS. 1A - 1D. The irradiated retinal spot on the fovea centralis **123** of the retina **122** is, as herein described, the origin of the re-emitted wavefront **130** measured by the sensor **356**. The output beam, probe laser beam **350** from diode laser **390** reaches the patient eye **120** by transmission through a linear polarizer and attenuator **392**, lens **394**, shutter **396**, and reflection off mirror **398** and in the polarizing beam splitter **378**. All these elements are fixed in position.

In one embodiment of the present invention, output of the diode laser **390** is essentially collimated and is focused onto a corneal surface of the eye **120** by lens **394**. As described in Application Serial Number 09/274,672 filed on March 24, 1999 for "Apparatus And method For measuring Vision Defects Of a Human Eye," and herein incorporated by reference, the projected probe laser beam **350**, collimated light from the diode laser **390**, is directed by a long focal length lens **394** for focusing on the anterior surface of the cornea **126** of the eye **120**, as illustrated by way of example with reference again to FIG. 1B, passing through the pupil and lens **124** of the eye **120**, and onto the retina **122** as a small measurable spot on the fovea centralis **123**. In one embodiment, the lens **394** comprises a zoom lens for varying the focus and moving the focus location as desired. By focusing on the cornea **126**, the measurement is

minimally dependent on the curvature of the cornea. However, other locations proximate the corneal surface are acceptable.

While diffraction and various aberrations are present, the present invention avoids the aberration effects from the cornea which typically dominate. The lens of the eye **120** contributes a relatively small aberration effect when compared to that of the cornea **126**. Further, and with regard to the selection of the lens **394**, selecting a lens with a short focal length would provide a relatively large incident angle of the beam **350**, a well focused point on the surface of the cornea **126**, and less aberration effects from the cornea. A small incident angle provides a larger focus point on the cornea **126**, but a more desirable smaller spot on the retina **122**, which spot size will depend on the wavelength and starting point size and focal length of the lens **394** selected. Embodiments of the present invention including lenses of approximate one half meter and 100 mm, by way of example, have been effectively used.

The polarizer **392** linearly polarizes the probe beam **350** into an s-state (by way of example, out of the plane of the drawing of FIG. 12). The angled interior interface of the polarizing beam splitter **378** reflects s-polarized light, so that light entering the eye **120** is s-polarized. A linear polarizer **400** is angled with respect to the polarizer **392** and works in conjunction with the attenuator to attenuate probe beam power delivered to the eye **120** to less than 10  $\mu\text{W}$ , by way of example. The diode laser **390** is triggered by an external electrical trigger signal **402**. A nominal illumination duration for eye measurement is 700 ms. The shutter **396** is included as an additional safeguard against overexposure of the eye **120** to the probe laser beam **350**. The shutter **396** is normally closed and is opened by an independent electrical trigger signal **404** synchronized to the laser trigger signal **402**.

By way of example, one retinal exposure for each illumination by the probe beam is  $10 \mu\text{W} \times 0.7 \text{ s} = 7 \mu\text{J}$ . Up to 10 repeat measurements may be obtained during a single patient examination session. Such exposures are well within the safety limits defined in the American National Standard for Safe Use of Lasers (ANSI Z136.1-1993, American National Standards Institute, New York, NY). In that reference, the maximum

permissible exposure (MPE) for "intrabeam" viewing a laser beam in the 400-700 nm wavelength range and the  $18 \times 10^{-6}$  to 10 second pulse duration range is  $1.8 \cdot t^{3/4}$  mJ/cm<sup>2</sup>. ( $t$  is the pulse duration in seconds). A limiting aperture for the eye is identified as approximately 7 mm in diameter. As a result, an allowable single-pulse energy is  $0.6927 \cdot t^{3/4}$  mJ. For a single 0.7 second pulse the MPE is 530  $\mu$ J, almost two orders of magnitude larger than a delivered energy per pulse, for the apparatus herein described. An additional calculation is performed to assess the safety of the repetitive exposures. The relevant calculation in the Standard multiplies the single pulse MPE by  $n^{-1/4}$ , where  $n$  is the total number of pulses in the exposure duration  $T_{max}$ . For the apparatus of the present invention, the 10-pulse safety limit is  $530 \mu\text{J/pulse} \times 10^{-0.25} = 298 \mu\text{J/pulse}$ , still a factor of 40 larger than the actual pulse energy focused into the eye.

As illustrated with reference again to FIGS. 12 and 12D, the fourth optical path 352 conveys the wavefront 354, earlier identified by numeral 130 with reference to FIG. 1B, emerging from the eye 120 to the wavefront sensor 356, herein described using a Hartman-Shack sensor by way of example for the wavefront analysis. The wavefront 354 re-emitted by the eye 120 in response to the probe beam 350 irradiation is conveyed to a CCD camera 406 by transmission through the polarizing beam splitter 378, the first afocal relay stage 358 lens combination, the 50/50 beam splitter 376, a trial lens holder 408, the second afocal relay stage 360 lens combination, reflection off mirror 410, and transmission through microlens array 412, as earlier described with numeral 33 with reference to FIG. 6. With the exception of the changeable trial lens holder 408, illustrated with reference to FIGS. 12F and 12G, all these optical elements are fixed in place on the surface of the platform 306.

The polarizing beam splitter 378 transmits only linearly polarized light in a p-state. The radiation of the probe beam 350 reflected from the corneal surface of the eye 120 will retain the incident s-state polarization and will not be appreciably transmitted by the beam splitter 378. In contrast, light that has been scattered off the retina of eye 120, light forming the wavefront 354 of interest, will be largely depolarized.

The p-polarized fraction of this light will be transmitted by the beam splitter **378**. Thus the beam splitter **378** selectively suppresses the corneal surface reflection that could otherwise complicate the wavefront measurement. A wavefront originating at the corneal plane of eye **120** is transferred to a plane of the trial lens holder **408** with unity magnification. This plane of the trial lens holder **408** provides an intermediate pupil plane and is included for placing an ideal N-diopter lens **409**, see FIGS. 12F and 12G, at the trial lens plane to change the spherical curvature of the wavefront **354** by N-diopters, without altering other aberration content. The capability to reduce/remove the general wavefront curvature in a preselected manner significantly extends the dynamic range in wavefront measurement, without degrading the measurement accuracy. Trial lenses **409a - 409m**, by way of example and herein described of varying spherical powers, ranging from -16 diopters to +8 diopters in two-diopter increments, are mounted on a rotating wheel **407** of the holder **408**. The wheel's axis of rotation is parallel to but offset from the optical axis **342**. Turning the wheel places one of a plurality of preselected trial lenses at the trial lens plane. The wheel has precision mechanical detents that register the selected lens properly in the optical path.

A narrow band-pass optical filter is also placed at trial lens holder **408** location just anterior to the lens position. This filter has maximum transmission for 670 nm wavelength radiation (the probe beam wavelength), and a bandwidth of approximately 10 nm (full-width-half-maximum). This filter is used to reject spurious light (from the fixation target illumination, the eye illumination, and the like) from the wavefront path.

In one embodiment, as herein described by way of example, each of the lenses of the second afocal relay stage **360** consists of three lens elements, two meniscus lenses and an interposed achromatic doublet. However, they are not identical, and their combined action serves to magnify the passing wavefront **130**. The wavefront **354** at the trial lens holder **408** location is imaged onto the surface of the microlens array **412** with a magnification of 1.22. Magnification of the wavefront image by this defined factor of 1.22 reduces the wavefront slope at each point in the image plane by the same 1.22 factor. This extends the measurement dynamic range of the device, again without decreasing accuracy. In addition, this magnification distributes the wavefront **130** over



more elements, CCD cells **38** as earlier described with reference to FIG. 6, in the microlens array **412**, thus increasing the number of slope measurements provided by the wavefront sensor **356**. The mirror **410** is included to fit elements of the apparatus **300** within the dimensions of the platform **306**. In addition, the mirror **410** also allows optical alignment adjustment for the microlens array **412** and the CCD camera **406** combination. As earlier described, by way of example, with reference to FIGS. 3-6, the microlens array contains a square array of microlenses which divide the incident wavefront into a transverse array of secondary "wavelets." These wavelets are focused onto a detector surface of the CCD camera, which is positioned parallel to the microlens array and one focal length posterior thereto. The pattern of focused wavelets in the CCD image is used to calculate the shape of the incident wavefront.

As illustrated with reference again to FIGS. 12 and 12E, the calibration beam path **362** provides the collimated beam **364** to the Hartman Shack wavefront sensor **356**. Wavefront data for the collimated beam **364** is used as a reference in reconstructing the aberrated wavefront **354** from the real eye measurement. The source for the collimated reference beam **364** consists of a diode laser **414** coupled to a beam expander **416**. In one embodiment of the invention herein described, the diode laser **414** used for reference is identical to the diode laser **390** used for the probe beam path **348**. The collimated reference beam **364** is conveyed to the CCD camera **406** by transmission through polarizer/attenuator **418**, negative lens and aperture **420**, aperture and positive lens **422**, reflection off mirror **424**, transmission through aperture **426**, reflection in the polarizing beam splitter **378**, transmission through the first afocal relay stage **358**, the 50/50 beam splitter **376**, the trial lens holder **408**, the second afocal relay stage **360**, reflection off the mirror **410**, and finally transmission through the microlens array **412**. Except for trial lens holder **408**, all these optical elements may be fixed in position on the surface of the platform **306**.

The optical element of the polarizer and attenuator **418** contains two linear polarizers and a neutral density filter. The linear polarizer furthest from the diode laser **414** polarizes the laser radiation in the s-state for maximum reflection in the polarizing

beam splitter **378**. The linear polarizer closest to the diode laser **414** is partially "crossed" with respect to the polarizer **378** to attenuate the laser power. The neutral density filter further attenuates the beam, such that the laser power reaching the CCD Camera **406** is optimal for calibration of the sensor **356**. The negative lens and positive lens of elements **418**, **420** expand the diode laser output and form the collimated reference beam **364**. Intervening apertures of elements **418**, **420** transmit only the central portion of the expanding beam with the most uniform intensity. The mirror **424** is included to reduce the overall dimensions of the apparatus **300**. The aperture **426** is conjugate to the corneal plane, and is included so that the collimated reference beam **364** illuminates approximately the same area on the microlens array **412** as would the wavefront **354** re-emitted by a maximally dilated eye.

By way of illustration, optical components suitable for use with embodiments of the present invention herein described by way of example, are provided with reference to Table 1. An electrical component layout of the apparatus **300** is illustrated with reference to FIG. 13, wherein a dashed box **428** indicates the platform **306** with the heretofore described element carried thereon. Except for the computer monitor, keyboard, and mouse all other electrical components are located within the frame under the optical table. Switches in the diagram are all located on a front panel **430** of the electronics **324** for ease in operator/technician access, as described earlier with reference to FIG. 9. Electrical power from the clinical facility is drawn by an isolation transformer, which in turn supplies power to an uninterruptible power supply (UPS). The UPS delivers power to three power strips carried in the frame **308**. The host computer **326** has a self contained On/Off switch, as do the three power strips. One power strip **432** supplies power to the shutter controller **434**, which commands the probe laser shutter **396** through the signal **404**, two dual power supplies **436**, each capable of providing both 5 VDC and 9-15 VDC output, the host computer **326**, a computer monitor **438**, and a third power strip **440**. One dual power supply supplies 5 VDC power to the two patient illumination lamps **336**, and 9 VDC power to the target illumination lamp **368**. A second dual power supply supplies 5 VDC power to both

diode lasers **390, 414**. A user-accessible 3-position switch **442** allows the system operator/technician to provide power to either the probe laser **390** or the calibration laser **414**, with a center switch position being the "off" state.

A third power strip **444** supplies power to the CCD electronics controller **446**.  
5 The power strip **440** also supplies power to cooling fans **448** located on the platform and within the frame.

By way of example and for illustration purposes, operation of the apparatus **300** may generally proceed with the operator/technician first activating each of the electrical elements, with the CCD electronics controller **434** being last to be enabled. The  
10 operator then activates the calibration laser **414** via the 3-position switch **442**. The operator then instructs the computer **326** to acquire a calibration wavefront measurement. The computer **326** relays this command to the CCD controller electronics **446**, which activates the CCD camera **406** to take a predefined exposure. The CCD controller electronics **446** also sends trigger signals **402, 404** described  
15 earlier with reference to FIG. 12, to the probe laser **390** and the probe laser shutter **396**. However, since the probe laser **390** is not powered at this point, no probe beam **350** is delivered. Calibration CCD data are transferred to the CPU of the computer **326**, and stored for later analysis. The calibration laser **414** is switched off at the end of the calibration procedure.

20 The technician/operator then proceeds to patient measurements. The output switch **442** at the dual voltage power supply **436** is positioned to a probe laser setting. The probe laser **390** is now in a "ready" state awaiting an additional trigger signal to operate. The operator then positions the patient appropriately in the apparatus **300** as earlier described with reference to FIGS. 9-11, with the assistance of an image from the  
25 video camera **338** displayed on the computer monitor, by way of example. With the patient correctly situated, the operator instructs the computer **326** to obtain wavefront data, as earlier described with reference to FIGS. 2-7. The computer **326** relays appropriate commands to the CCD electronics controller **446**, which triggers the probe laser **396** to fire, triggers the shutter controller **434** to open the probe laser shutter **396**,

and exposes the CCD camera 406. CCD camera image data is transferred back to the computer 326. The computer 326 includes software that analyzes the patient and calibration data to calculate the patient wavefront profile for use in the corrective surgery, as earlier described with reference to FIG. 8, by way of example. At the end of the data collection, the operator shuts down the electronics, starting with the CCD electronics controller 446. The software integrated into the apparatus 300 may be described, by way of example, as including: A graphical user interface (GUI) to allow the technician to perform all desired operations to enter and save patient information and perform the desired measurements; database and file system interfaces to allow for the saving and tracking of patient information, measurement, and hardware details; control of the electro-optical and electro-mechanical components as necessary in order to be able to accurately and safely perform the desired measurements; and processing of the measurement data to generate mathematical descriptions of the aberrations (the optical path difference) measured in the subject eye. By way of example, these mathematical descriptions of the aberrations can then be used in a LADARVision® system to perform optimal refractive surgical procedures, which system is available from Autonomous Technology Corporation, a wholly owned subsidiary of Summit Technology, Inc.

By way of further example, patient measurement and apparatus configuration information is stored in multiple tables in a Microsoft Access™ 7.0 database. The interface to this within the code is based upon the Microsoft Foundation Classes (MFC) wrapper to the Microsoft Jet Engine. The framework generates a Structured Query Language (SQL) to create, retrieve and update records in the database. Use of the Microsoft Access application to access the data is not needed. In one embodiment of the present invention, the following data may be stored in the database: patient information - name, address, medical record number, and the like; measurement Information - geometry, time of measurement, and the like; and system Information - hardware serial numbers and key hardware parameters.

Additionally, the software may be developed with two operating levels - password-protected and not-password-protected. From within the password-protected-

mode, the technician/operator has access to system configuration information and features necessary for system setup and maintenance that are not accessible from the not-password-protected mode. All patient entry and measurement capabilities are available from the not-password-protected mode. All patient information desired in order to be able to uniquely identify and track the patient is entered via the graphical user interface (GUI) and stored in the Microsoft Access database. Selecting the "Patient Data" menu item brings up a patient information data information screen, from which the technician can enter new patient data as well as being able to review and edit existing information. The patient data that can be stored and retrieved, typically includes: name, address, medical record number, data of birth, phone number, sex, manifest and cycloplegic refractions and vertex distance as well as centration information.

Centration information that is measured via a centration process and stored as part of the patient record describes the position of the center of the constricted pupil with respect to the center of the limbus. This information is used in aligning the patient for the measurement where the goal is to align the visual axis of the eye with the optical axis **342** of the apparatus **300**. When the centration procedure is invoked a list is displayed of all patients that have been entered into a database operable with the apparatus **300** but have not yet had the centration steps performed. The monitor displays all patient information including a review of centration information, or alternatively, for just those patients entered for a given time period. In order to perform centration for a given patient and eye, that patient is selected from this list by clicking on the desired patient/eye with the mouse. An example of the centration process is illustrated with reference to FIG. 14. Once a patient has been selected, the patient is instructed to look into the apparatus **300** and at the fixation target **366**, as earlier described with reference to FIG. 12.

By way of further disclosure, the fixation target **366** is, as earlier described, included so that the patient **302** can stare along the optical axis **342** of the apparatus **300**. For best fixation, the target should be clearly visible to the patient. However, care should be taken to see that the patient does not attempt to accommodate when fixating

on the target. This would occur if the target were optically closer than the patient's infinity focal plane. If the patient did accommodate, i.e., if the lens in the eye changed shape to provide increased focusing, then the eye would appear excessively myopic during the wavefront measurement. To avoid this, the fixation target optics are  
5 adjusted so that the target appears to lie optically just beyond the patient's far-field focus. Thus for each patient the target will appear relatively clear, but not in sharp focus. The patient may initially try to accommodate to improve the sharpness of the image, but will eventually find that the clearest image is seen for the most relaxed (non-accommodative) state. This technique is known as "fogging," and is routinely  
10 performed by optometrists when doing clinical evaluations. The eye drops used to dilate the eye for the measurement also reduce the lens' ability to accommodate, thereby further ensuring valid wavefront measurement.

With reference again to FIG. 14, an image of the patients eye **120** is frozen. Two  
15 reticles **450**, **452** are then used to locate the centers **454**, **456** of the constricted pupil and limbus, respectively. Each reticle **450**, **452** can be moved and sized - one reticle **450** is positioned over the perimeter of the constricted pupil **458** and the other reticle **452** over the limbus **460**. Once they have been correctly located the information is saved to the database. This can be performed for as many patients as is desired and the centration procedure is then exited.

20 For illustration, a sequence of events followed in measuring the refractive errors in an eye and computing the corresponding optical path difference (OPD) is illustrated with reference to FIG. 15. By way of example, steps include performing a reference measurement **462**. To provide a reference with which to compare the measurement of the eye **120** and also to check the alignment of the apparatus **300**, a reference  
25 measurement is made using the collimated laser light **364**, as earlier described with reference to FIG. 12. The software forces the operator to make at least one such measurement at the start of each day and an additional one at the end of each day. More reference measurements can be performed as desired by the operator. When patient measurements are performed, the measurement records in the database  
30 identify which reference measurements correspond to each measurement, i.e., which

reference image was the latest one done prior to the measurement. A "Perform Reference Measurement" screen may be provided for viewing a sample reference image.

5 A next step includes selecting a patient and eye to measure **464**. The patient and eye to be measured may be selected from a "Patient Select" dialog screen. It is desired that all patients are displayed along with a check mark to show whether or not centration has been performed for that patient. If a patient is selected that has not yet had centration performed then the operator is informed of this and no measurement can be performed. Once a valid patient/eye has been selected to be measured then the perform measurement dialog is displayed which includes GUI buttons necessary in order for the operator to perform and check the measurement.

10 A next step includes aligning the eye using the video camera and reticles **466**. The apparatus **300** is operated with the visual axis of the eye aligned, as close as is practically possible, to the optical axis **342** before performing a measurement. The center of the constricted pupil **454** is used as the approximate anatomical landmark for the visual axis. Given that the eye **120** is dilated when the measurement is performed, it is not possible to directly determine this center. However, the centration procedure performed on each patient defines the center of the constricted pupil **454** with respect to the limbus **460** and thus it is possible to use the position of the limbus to place the eye **120** in a desired location.

15 As illustrated with reference to FIG. 16, a reticle **468** is displayed on screen that is offset from the optical axis by the appropriate amount such that when the limbus **460** of the eye **120** is aligned to this reticle **468**, the eye **120** is positioned as desired. Prior to taking the measurement, it is the operator's responsibility to ensure that the patient is positioned appropriately such that the limbus **460** is aligned with the reticle **468** while the patient is looking at the fixation target **366**.

20 A measurement is then performed **470**. Once the eye **120** is correctly aligned, the operator presses an "acquire" button to perform the wavefront measurement of the patients eye. The system response to the acquire command is as follows:

- 30 1. Video image is frozen

2. Probe beam laser is activated
3. External shutter is opened so that the probe beam can reach the eye
4. CCD shutter opens and the CCD is exposed to the re-emitted wavefront (1-4 generally performed simultaneously)
5. CCD shutter closes and the exposure is completed
6. CCD data is transmitted to the computer
7. External shutter closes and the probe beam turns off.

The software continually checks the status of the CCD electronics and the temperature of the camera and only allows measurements to be taken when everything is working nominally.

A review of eye and apparatus geometry is accepted or rejected **472**. Although it is not necessary for the eye **120** to be perfectly aligned with respect to the optical axis **342** (the software compensates for minor misalignments), it is desirable for it to be close. The eye **120** will have been aligned prior to the measurement but uncontrollable eye motion (e.g. saccades and loss-of-fixation) may make the alignment sub-optimal at the time of the exposure. To check that the alignment is acceptable, the video image of the eye is frozen at the time the measurement is taken. The operator then aligns a reticle to the limbus ring and presses a "check geometry" button on the GUI. If the software determines that the alignment is not acceptable, the operator is informed of this and a new exposure is made as desired. By way of example, and with reference to FIG. 17, optimal measurement as herein described, would have the limbus **460** aligned to circle B. In actuality, the eye **120** was offset during the exposure and the limbus **460** was aligned to circle A. The difference between these two states is shown by the line A'B'. The software determines whether or not the image is acceptable based on the length of A'B'.

It is also at this point that the operator records the rotational state of the eye. Prior to the wavefront measurement, a pattern of four line segments **474** arranged in an "X" pattern **476**, as illustrated with reference to FIG. 18, around the periphery of the cornea are applied to the eye using a mechanical instrument. The pattern **476** consists of two pairs of collinear line segments **474** angled at 45° with respect to each other.



Each line segment **474** is 4 mm long, and collinear segments are separated by 7 mm. At the same time the limbus ring reticle is aligned with the actual limbus in the frozen video image, an X reticle that matches this pattern is aligned to the applied eye marks in the frozen image. The orientation information is then saved by the software along with the limbus position data.

As a next step in the process, the CCD image is processed, accepted and saved, or rejected **478**, as illustrated with reference again to FIG. 15. If the geometry of the measurement is acceptable, it is then probable that the quality of the CCD image will be high. It is desirable, however, to check that this is so. The software processes the image and then presents an auto-scaled image to the operator to review. If the software determines that the image is unacceptable then the operator will be informed of this and a new exposure made. If the user decides that the image is unacceptable for whatever reason then the image can be manually rejected at this stage. An example of an unacceptable image is illustrated with reference to FIG. 19. In this example, a significant portion **479** of the image is obscured in some manner, resulting in wavefront data for only part of the pupil. By unacceptable, it is meant that such an image is not believed to result in the accuracy and precision of measurement that is desired for surgical procedures which are obtained by the present invention. It does not mean that such as illustrated may not be usable in any sense.

Once a valid measurement has been made the next step **480** is to measure the local slopes of the wavefront **130**, as earlier described with reference to equations herein presented. As described with reference to FIGS. 4-6, it is necessary for the software to compute the centroids **116** of the clusters of light on the CCD array **38** and then determine the distances of each of these centroids **116** from the corresponding reference centroids **29**. The centroids are determined by first computing which pixels should be processed and grouping them together into clusters. The intensity-weighted centroid of each cluster is then computed. As illustrated with reference to FIG. 20, an example of an image from a myopic eye with the computed centroids **482** of cluster **484** marked by "X"s is shown. FIG. 21 illustrates a close-up of one of the clusters **484** and displays not only the centroid **482** but also the pixels **486** used in the centroiding

calculation for the cluster 484. CCD pixels 488 processed in the centroiding algorithm are marked by dots. This algorithm, by way of example, isolates centroids by use of a spacial filter which removes stray light signals that create noise for the CCD image. Such filtering may be desirable before calculation of light cluster positions.

Without filtering, computation of the cluster centroids may be made difficult as a result of noise on the image such that individual pixels with no actual data content may be brighter than pixels containing relevant data, speckle in the image may result in valid data clusters having irregular profiles with significant variation in intensity of adjacent pixels, haze or background noise may be high relative to the actual data or may be non-uniform across the image, intensity of valid data may be non-uniform across the image, scatter from different parts of the eye may result in spurious signals on the image, and high aberrations in the eye may significantly distort the clusters of valid data, by way of examples. The spatial filter permits a re-computation of the brightness of each pixel in a bitmap using a weighted averaging technique that considers surrounding pixels. In a particular application herein described for illustration and by way of example, the spatial filter is designed to yield a maximum value when centered on valid data, reduce an effect of individual bright pixels or small groups thereof, normalize background levels, smooth valid data profiles, and simplify the task of extracting the valid data from background noise or haze. One filter employed in one embodiment of the present invention is square ( $n \times n$ ) and includes real values (positive and negative) assigned to each pixel. The filter is designed to be optimally matched to images obtained from eyes with high, yet measurable, levels of aberration. By wave example, a cross-section through the filter is illustrated with reference to FIG. 23A. An effect of applying such a filter improves an image such as illustrated with reference to FIG. 23B to the image illustrated with reference again to FIG. 20, by way of example, a cleaner image and one that is easily processed for identification and computation of cluster centroids. By applying the filter, images that would otherwise be deemed to noisy or of insufficient quality to process, can now be processed and desired wavefront information computed.

The center of each centroid is calculated using a standard center of mass algorithm based on light intensity. The clusters and centroids illustrated with reference

to FIG. 22 are illustrated with the locations of the corresponding reference centroids **490** also visible. The open circles in this figure indicate the locations of the reference centroids. Lines connect these with the associated sample centroids **482**. From the distances between the reference and measurement centroids **490**, **482** respectively, and the distance between the lens array **33** and the CCD plane **36**, described with reference to FIG. 6, the local slopes are calculated. Given these local slopes and information about the apparatus setup, including any and all magnification factors, it is then possible to determine the local slopes at the pupil plane and, from these, and compute the optical path difference of the eye being measured.

A description of the wavefront is then made **492**. As earlier described, the reconstructed wavefront is described via a set of Zernike polynomials. The number of locations on the eye **120** at which the local slopes are determined (i.e. the number of sample points) greatly exceeds the number of terms in the polynomials that will describe the wavefront. A least-squares-fit calculation is done to find the coefficients that best describe the surface. The order of the polynomial used is sufficient to describe not only the spherical and cylindrical refractive powers (2<sup>nd</sup> order) but also the levels of coma (3<sup>rd</sup> order) and spherical aberration (4<sup>th</sup> order) present.

An example of the computed Zernike coefficients for an eye and the corresponding wavefront reconstruction **493** is illustrated with reference to FIG. 24A. By way of example, for the wavefront illustrated with reference to FIG. 24A, the spherical and cylindrical powers computed from the wavefront are -1.60/-1.13 x 150.4. The corresponding values obtained by an optometrist performing a phoropter examination (converted to the corneal plane) were -1.47/-1.19 x 150. The standard measurements of spherical and cylindrical powers agree well with the computation of spherical and cylindrical powers, but there are also higher order aberrations present. By way of further example, FIG. 24B illustrates just these higher order aberrations **495** on the same scale as the plot of FIG. 24A.

With regard to the optical path difference (OPD), scaling an optical path difference profile,  $OPD(x,y)$ , by a refractive index difference (cornea to air) is not the only step included to calculate the correct ablation profile. In addition, the present

invention allows for a treatment on the curved corneal surface, while the wavefront measurement was made at a plane tangent to the cornea, as illustrated with reference to FIG. 25, which is exaggerated to illustrate the effect. The image plane of the wavefront path is the lenslet array plate. The object plane of the wavefront path is the reference plane 494. In this highly-exaggerated myopic case, herein described by way of example, one light ray 496 emerging from the eye 120 at transverse location  $a$  is detected at a transverse location  $b$ . The wavefront reconstructed from sensor data will have the slope of this ray at location  $b$ . Although this is true of the wavefront at the reference plane 494, simple scaling of this wavefront would yield an ablative treatment at corneal location  $b$  that may not be entirely correct. In actuality this effect is small. The radius of curvature of the cornea is typically on the order of 7.5 mm. (a range of 7-8 mm encompasses most eyes.) At a transverse location 3 mm from the corneal apex, the distance from the corneal surface to the reference plane is only  $\sim 0.63 \mu\text{m}$ . For a 10 diopter myope, a light ray exiting the cornea at  $a = 3.0 \text{ mm}$  will cross the reference plane at  $b = 2.98 \text{ mm}$ . The difference between  $a$  and  $b$  in this example is only  $20 \mu\text{m}$ . Although small this geometric effect is systematic, having progressively greater impact on the measurement with increasing radial distance from the corneal apex. To increase the accuracy of the treatment profile, compensating for the curved geometry may be performed in the following manner:

1. Wavefront slopes are calculated at each measurement point in the reference plane.
2. The cornea is assumed to have a nominal radius of curvature ( $\sim 7.5 \text{ mm}$ ).
3. The wavefront slopes measured at the reference plane is projected back onto the nominally curved cornea. The wavefront is measured to have a certain slope at  $b$  in the reference plane, described above. It is a straightforward mathematical process to calculate the point  $a$  where this ray exited the cornea.
4. The wavefront is reconstructed based on the measured slopes at the calculated corneal locations. This wavefront is used in determining the ablation profile.

As above described, a wavefront measurement has the patient correctly positioned at the apparatus 300. The eye 120 being measured is at the correct location and looking in the appropriate direction. Based on analysis of the allowable eye-positioning tolerances, the apparatus 300 of this embodiment of the present invention provides the following patient position information:

The capability for ensuring that the subject eye is at the right location along the longitudinal (z) axis of the apparatus with an accuracy of +/- 1 mm.

The capability for ensuring that the subject eye is correctly positioned laterally with respect to the apparatus (i.e., in x-y) with an accuracy of +/- 1 mm.

The capability for ensuring that the subject eye is correctly positioned in angle with respect to the apparatus (i.e., the difference between the visual axis and the optical axis of the system) with an accuracy of +/- 0.5 degrees.

The capability for aligning an on-screen reticule to a set of marks applied to the eye outside the limbus to record the rotational orientation of the eye (i.e., about z) with respect to the apparatus with an accuracy of +/- one degree.

Once in position, the patient's eye can be successfully examined by the wavefront sensing technique. This embodiment of the apparatus includes a sufficient dynamic range to measure eyes over the expected scope of refractive errors. In addition, the apparatus detects complex aberrations, and does so with sufficient accuracy to serve as the basis for ablative treatment.

The following list provides range and accuracy parameters, by way of example, for clinical wavefront measurements that can be obtained by this embodiment of the apparatus. This list is provided by way of illustration and does not limit the scope of the present invention.

1. capable of measuring wavefronts with spherical refractive powers in the range +6 to -15 diopters and cylindrical powers in the range 0 to -6 diopters.
2. capable of measuring coma and spherical aberration.
3. capable of measuring refractive errors over a pupil zone of up to 8 mm in diameter.

4. able to measure the refractive errors within the specified ranges to an accuracy of 0.042  $\mu\text{m}$  RMS in air.

A computation of a shot pattern is performed in the LADARVision® system. The Zernike coefficients computed in the manner described here are imported into the LADARVision® system along with all other d measurement and patient information and used along with LADARVision® system parameters to compute the optimal number and placement of shots.

One embodiment of the present invention for a calculation of a treatment laser spot pattern includes an ablation effectiveness distribution over the corneal surface. One embodiment of the present invention, as herein described, optimizes refractive surgery ablation profiles so that post operative aberrations are minimized. One treatment profile takes into account information beyond just that of pre-operative aberrations. As the reader will appreciate, the use of wave front measurement devices has provided greater insight into the effectiveness of current excimer ablation profiles. Analysis of multiple patients for pre and post laser reflective surgery has resulted in a model for describing an effectiveness of a laser ablation as a radially symmetric attenuation function. One embodiment of the present venture provides for this attenuation function. As illustrated by way of example with reference to FIGS. 26A and 26B, a difference exists between an intended change in corneal depth using laser ablation, and an achieved change. FIG. 26A illustrates an intended and achieved profile for surgery on a myopic eye, while the 26B illustrates an intended and achieved profile for surgery on a hyperopic eye. The ablation depth versus normalized radial profile plots of FIGS. 26A and 26B are representative of multiple surgeries analyzed. A constant attenuation independent of radial position results. Sometimes the attenuation is zero. In addition, a radially symmetric attenuation function results. Such a function can be described by an equation of the form:  $\text{Ablation Efficiency}(p) = A\{1 + Bp^2 + Cp^2\}$ , where  $p$  is a normalized radial position, and A, B, and C are coefficients describing the attenuation function. The attenuation function may be graphically described, by way of example, with reference to FIG. 26C. As a result, an embodiment of the present invention takes a previously unknown efficiency or attenuation function and modifies

treatment profiles accordingly so that a desired outcome is achieved. By way of illustration and example, this may be accomplished by taking a desired change in corneal depth (e.g. a nominal ablation profile), and dividing the nominal profile the attenuation function. This yields a new profile which, when ablation is performed, will result in the desired profile. One approach is to compute the Zernike description of the ablation profile as earlier described, and divide the resulting Zernike polynomial by the attenuation function to compute a modified Zernike description of the ablation profile that is to be used with the ablation laser system. By way of example, if the  $P_{\text{DESIRED}}$  is the desired change in corneal depth (i.e. the desired achieved ablation profile) and  $P_{\text{INPUT}}$  is the profile to be entered into the ablation laser system, then  $P_{\text{INPUT}}$  may be defined by:

$$P_{\text{INPUT}}(\rho, \theta) = P_{\text{DESIRED}} \div A\{1+B\rho^2+C\rho^4\}$$

With reference again to FIG. 6, and by way of further example, the output from wavefront analyzer 26, e.g., the Zernike expansion of equation (19), can be used in a variety of ways. For example, the output may be used to continually or periodically monitor the progress or effects of an ophthalmic procedure, with such stored on disc or transmitted via e-mail, and the like. In addition, the measurement of the eye and the resulting surgery need not take place at the same site. The output could also be used to develop an optical correction for the eye 120. The optical correction will make the aberrated wavefront 130 appear approximately as the planar wavefront 110. As described above, the optical correction can be implemented in a variety of ways. In each case, the output of the wavefront analyzer 26 is input to a processor 90 which converts the Zernike expansion of equation (19) into a form suitable for being implemented as one of the possible optical corrections. Alternatively, the processor 90 may also be implemented at the processor 40 of the wavefront analyzer 26, described earlier with reference to FIG. 6.

By way of further example, the processor 90 can be used with preselected Zernike coefficients from the expansion of equation (19) to generate a standard spherocylindrical correction for a lens grinder 92 to produce a convectional optical lens, e.g., a lens for glasses, a contact lens, and the like.

In one embodiment of the present invention, herein presented by way of example, the processor **90** includes a modification of the Zernike reconstruction of the aberrated wavefront **130** by the index of refraction of the cornea **126** minus that of air, having value of 1, as earlier described, to calculate an amount of corneal material to be ablated at each corresponding (x,y) location on the cornea **126**. This information regarding the amount of corneal material can be used in conjunction with a laser beam delivery system **94** that typically has eye tracking capability. The laser beam delivery system **94** including the eye tracker is placed in line with the optical axis of the apparatus **11**, as illustrated again with reference to FIG. 7. The eye tracker portion allows the apparatus **11** to respond to unwanted eye motion. The system **94** would typically focus short pulses or "shots" of ablating laser light onto the cornea **126** to remove a specified thickness  $t$  of material at each location. This is shown diagrammatically in FIG. 8 where the uncorrected surface of the cornea **126** is referenced by numeral **126A** and the corrected surface of cornea **126** after ablation is referenced by numeral **126B**. In accordance with the present invention, the ablation thickness  $t$  is specified across the aperture of the cornea measured, e.g., the 6 millimeter circle to which the eye's pupil was dilated during the measurement of the eye. Outside the prescribed treatment circle, a tapering blend zone of partial ablation may be added to minimize severe changes in corneal curvature and hence lessen regression. The laser beam delivery system **94** removes thickness  $t$  to achieve the optical correction, which results in the corrected cornea surface **126B**. Note that the optical correction is not concerned with the ultimate corneal topography, but instead removes corneal material to achieve an optical correction that takes into account all ocular aberrations of the eye **120**. This is important because the shape of the corneal surface can be independent of the correction  $d$  because the eye's vision depends on numerous factors besides corneal curvature. Hence, the best corneal surface topography for optimal vision may be far from regular in that it may compensate for the errors in the eye's other surfaces. Thus, it is apparent that the present invention can be used to provide corneal surface corrections other than the conventional spherical and/or cylindrical corrections.



As described earlier with reference to FIG. 12, the apparatus **300** of the present venture includes first and second afocal relays stages **358**, **360**. To retain the benefit of wavefront magnification, as a means of increasing the dynamic range of the wavefront sensor **356** to accommodate patients with large refractive errors, while at the same time allowing for incorporation of a small format, inexpensive camera to record the wavefront slope data, a modification **500** to the apparatus **300** as illustrated with reference to FIG. 27A is provided.

By way of example, a lens array may also be positioned and configured as illustrated with reference to FIG. 27B, wherein a portion of the apparatus **300** of FIG. 12 includes the first and second afocal stages **358**, **360** within the optical axis **342**, and the wavefront sensor **356** consist of the microlens array and CCD camera separated by a fixed distant, as earlier described with reference to FIG. 6. This optical path through the afocal relay stages results in an image of the corneal plane **502** at the lenslet array, i.e. at the entrance face of the actual wave front sensor **356**. This can be accomplished by a single afocal stage. As earlier described with reference to FIG. 12, the apparatus **300** includes an intermediate image plane as insertion point, the holder **408**, for a trial lens. Placing a spherical lens into the optical axis **342** at the first image plane, in theory, could be used to remove the defocus wavefront error. This would potentially expand the dynamic range of the apparatus **300**. However, the trial lens approach is a moving mechanism that can position lenses at the first image plane with tremendous accuracy in repeatability. It is highly desirable that alternative means be developed to address dynamic range.

One way to accomplish this is to magnify the corneal plane image at the lenslet array with the afocal stage **360**, earlier described. Magnification of the wavefront reduces the wavefront slope, so that the displacement of the focused lights spots on the CCD is decreased. The chosen magnification factor used with the apparatus **300** second afocal stage **360** is approximately 1.2 which is sufficient to cover the desired range in refractive errors. A magnification factor in excess of 1.5 is desirable for expanding the use of the apparatus **300**. However, simply magnifying the corneal plane

has a drawback in that it necessitate a large aperture wavefront sensor. That is, both the lens array and the CCD camera preferably have large cross-sectional areas to encompass the magnified image of the point of plane. This is not a significant issue for the lens array. However, a large format CCD camera is quite expensive and such cameras are only available from a limited number of vendors.

To resolve such concerns, the modification 500 illustrated with reference again to FIG. 27A is provided. The corneal plane 502 is imaged at a reference plane 504 by an afocal relay stage 506, which magnifies the corneal plane by a preselected amount. The lenslet array 412 is placed at the reference plane 504. Focused spots of light from the eye 120 are produced at the lenslet array focal plane 504. Rather than place the CCD detector face at the reference plane 504, an optical train 508 is inserted to image the array focal plane 413 at yet another plane, a final image plane 510, at which plane the CCD detector face is positioned. The afocal relay stages 358, 360 described earlier with reference to FIGS. 12 and 27B, may or may not be included, as desired. However, the magnification of the array focal plane at the final image plane 510 is provided. This allows a small, relatively inexpensive, active area camera to be used as the light recording element in the wavefront sensor. Details of optical design including magnification specifics can be adjusted to maximize performance for a given camera and lens array plate specification.

The advantages of the present invention are numerous. A totally objective approach is presented for measuring ocular aberrations. The approach is effective for a wide range of vision defects. Accordingly, the present invention will be of great utility in a wide variety of clinical applications. For example, the calculated Zernike coefficients can be used to develop a completely objective lens prescription or a corneal correction that could be accomplished with laser ablation. In addition, each of the wavefront sensor embodiments provides for a greater degree of accuracy over the prior art with respect to measuring wavefront deflections. Further, the present wavefront sensor can be adjusted in terms of gain simply by adjusting the separation distance between the imaging plane of the sensor and the planar array of light-sensitive cells.

The objective measurement of the present invention will also find great utility for a large variety of applications where the "patient" is unable to provide feedback as d by conventional eye diagnosis. For example, the present invention could be used to evaluate the eyes of any patient not possessed of demonstrative communicative skills, e.g., babies, animals, dead specimens, as well as any constructed optical system, since the present invention is an objective analysis not requiring any assessment from the "subject". All that is necessary is for the subject's eye to be properly positioned so that proper optical access to the eye can be obtained.

The present invention will also be used in the area of identification should it be determined that each eye's Zernike coefficients are unique. Then, the present invention would find great utility in the fields of law enforcement, credit card/bank security, or any other field where positive identification would be beneficial.

Although the invention has been described relative to a specific embodiment thereof, there are numerous variations and modifications that will be readily apparent to those skilled in the art in light of the above teachings. It is therefore to be understood that, within the scope of the appended claims, the invention may be practiced other than as specifically described.

1           **83.**    The method of claim 57 in which the vision of the eye is enhanced from  
2 an eye requiring a correction of greater than 8 diopters to an eye having better than  
3 20/40 vision.

1           **84.**    The method of claim 57 in which the vision of the eye is enhanced from  
2 an eye requiring a correction of greater than 8 diopters to an eye having at least 20/20  
3 vision.

1           **85.**    A method for enhancing vision in an eye, the method comprising:  
2                   determining an optical path difference between a plane wave and a  
3 wavefront emanating from an eye;  
4                   producing a plurality of laser beam shots;  
5                   applying said plurality of laser beam shots to the eye in a manner to  
6 create two different focus zones and that is based in part on the optical path difference  
7 between the plane wave and the wavefront emanating from the eye; and,  
8                   said plurality of laser beam shots removing tissue from the eye in a  
9 manner that reduces the optical path difference between the plane wave and the  
10 wavefront emanating from the eye; whereby the vision of the eye is improved.